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source
1. 803
/organism="Homo sapiens"
/clone_id="3906"
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/notes="Creation of genome-wide Protein Expression Libraries using Random Activation of Gene Expression", Nature Biotechnology, in press. Note that even though the cell type indicated is H1080, since a random activation method was used, these sequence tags are not necessarily expected to be in 171 g of normal circumstances."
BASE COUNT      222 a      171 g      238 t
ORIGIN
Query Match      95.9%; Score 701.8; DB 12; Length 803;
Best Local Similarity 58.2%; Pied. No. 6.9e-188;
Matches 720; Conservative 0; Mismatches 12; Indels 1; Gaps 1;
Oy 1 CAAATCGGAAAAGATCCCTTTCCTCCATGCTGCTGATATTTAAATGCA 60
Db 761 CAAATCGGAAAAGATCTGTTTCTCAACCTGTGGCAATTAATTTAAATGCA 702
Oy 61 GCGAGAGGAGGATGCTACTTATTCATTCATTCAGACAGACGCTGGAAAGACCTCTCC 120
Db 701 GCGAGAGGAGGATGCTACTTATTCATTCATTCAGACAGACGCTGGAAAGACCTCTCC 642
Oy 121 -AAGATATTTATCTTATATGGAACATTTGGCCGAGACACAGAGTGAACAGAG 179
Db 641 AAGGATATTTATCTTATATGGAACATTTGGCCGAGACACAGAGTGAACAGAG 582
Oy 180 CCGTCCAGACCAAGTGAATGCTGAGAACCTTCATATGTCATCTGTAAGTGTGA 239
Db 581 CCGTCCAGACCAAGTGAATGCTGAGAACCTTCATATGTCATCTGTAAGTGTGA 522
Oy 240 TCGTTGAGATGTCCTCTTCATCATCGTGGCCATCTGCTGAGCACTGTGAATTCAG 299
Db 521 TCGTTGAGATGTCCTCTTCATCATCGTGGCCATCTGCTGAGCACTGTGAATTCAG 462
Oy 300 GCGGAGGAGACATGAGAGCCCTACACACCATATCTGTAAGAGCTGCGACGAAAGT 359
Db 461 GACGGGAGACATGAGAGCCCTACACACCATATCTGTAAGAGCTGCGACGAAAGT 402
Oy 360 ACAAGAGCCCAATCTGAATCTGAAGAAATCGAAGGCGACATCATGAGAAATCTGGT 419
Db 401 ACAAGAGCCCAATCTGAATCTGAAGAAATCGAAGGCGACATCATGAGAAATCTGGT 342
Oy 420 GCGTGGGCTCAAAATGTCCTGCTGAGAGAGCAAGCCACCAAGCTGAAATCTGAGT 479
Db 341 GCGTGGGCTCAAAATGTCCTGCTGAGAGAGCAAGCCACCAAGCTGAAATCTGAGT 282
Oy 480 CCGAGATCGAAGAGATCCCATGCGACGAGGAGAACCAATGCTGCTTAAAGAA 539
Db 281 CCGAGATCGAAGAGATCCCATGCGACGAGGAGAACCAATGCTGCTTAAAGAA 222
Oy 540 AGTATATCTCTGCTCTGTTGTGAATTCATGAGAAATTTATGTTGGTGGCAATAG 599
Db 221 AGTATATCTCTGCTCTGTTGTGAATTCATGAGAAATTTATGTTGGTGGCAATAG 162
Oy 600 ATATGATGACATTCATCTCACTGATTTATGCTGCTGTGAGCAAAATTTTCTGCTG 659
Db 161 ATATGATGACATTCATCTCACTGATTTATGCTGCTGTGAGCAAAATTTTCTGCTG 102
Oy 660 AAGACCTCTTTTCTTCCGCGCAAGGAAATGATGCTGTTGAGCAATTAATGATGAAA 719
Db 101 AAGACCTCTTTTCTTCCGCGCAAGGAAATGATGCTTATTAATCAATCATGATGAAA 42
Oy 720 TAAAGCAAAATTT 732
Db 41 TAGAGCCAAATTT 29

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LOCUS      A1654552      429 bp      mRNA      linear      EST 17-DEC-1999
DEFINITION      w46b12.x1 NCI-CGAP GC5 Homo sapiens cDNA clone IMAGE:2308895 3
similar to SW:NRK_HUMAN P15872 IJR SLOW VOLUME-GATED POTASSIUM
A1654552 PROTEIN 1; mRNA sequence.
ACCESSION      A1654552
VERSION
KEYWORDS
SOURCE
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
AUTHORS      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap
TITLE      National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
JOURNAL      Tumor Gene Index
COMMENT      unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgap@fremail.nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
Bonaldo, Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
clone distribution: NCI-CGAP clone distribution information can be
found at: http://www.ncbi.nlm.nih.gov/ncicgap/ncicgap.html
Insert length: 771. Std Error: 0.00
Seq primer: -40bp from Gibco
High quality sequence ntbp: 411.
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/tissue="pooled germ cell tumors"
/lab_host="DH10B"
/notes="Vector: pUT7D-pec (pharmacia) with a modified
polylinker; Site 1: Not I; Site 2: Eco RI; plasmid DNA
as circles were made in vitro. Following Hsp
hybridization
this DNA was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from a pool
of 5,000 clones made from the same library (cloneids
1257096-1258631, 1459064-1470993, and 1475592-1476743).
Subtraction by Bento Soares and M. Fatima Bonaldo.
BASE COUNT      127 a      100 c      104 t
ORIGIN
Query Match      56.3%; Score 412.2; DB 9; Length 429;
Best Local Similarity 99.0%; Pied. No. 7.1e-106;
Matches 411; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
Oy 110 GAGCTCTTCGAGAGATTTATTAATCTATATGAGCAATTCGCGCCAGCAACAAAGCT 169
Db 12 GAGCTCTTCGAGAGATTTATTAATCTATATGAGCAATTCGCGCCAGCAACAAAGCT 71
Oy 170 GAGGAGAGGCGCTTCAGACCAATTTGATCTGAGAAATCTCTCAAGTCATCCGTGAC 229
Db 72 GAGGAGAGGCGCTTCAGACCAATTTGATCTGAGAAATCTCTCAAGTCATCCGTGAC 131
Oy 230 CTGATGATGATATGGAATGCTCTTTCATCATACATGCTGGACACCTGCTGAGACCTGTG 289
Db 132 CTGATGATGATATGGAATGCTCTTTCATCATACATGCTGGACACCTGCTGAGACCTGTG 191
Oy 290 AATATCCAGAGAGGAGGAGACTCCATACAGACCCCTACACAGCATATCTGTGAGAGGATG 349
Db 192 AATATCCAGAGAGGAGGAGACTCCATACAGACCCCTACACAGCATATCTGTGAGAGGATG 251
Oy 350 CAGGAAAGTACAGAGGCCAATCTTGATCTGAAAGATGACAGGCCATCTCATGAG 409
Db 252 CAGGAAAGTACAGAGGCCAATCTTGATCTGAAAGATGACAGGCCATCTCATGAG 311

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OY 410 AACATGTCGCGCTGGCTTCAAAATGTCCTCCCTGATAGAGGAAAGGACCAACACTTA 469  
 DB 312 AACATTTGGGCGCGCTGGCTTCAAAATGTCCTCCCTGATAGAGGAAAGGACCAACACTTA 371  
 OY 470 CACGACGACGTCGAGACATGAGATGCGGCGGACGAGGCGCAATTCGAAATTCGCTT 527  
 DB 372 CACATGCGTCGACGACATGAGATGCGGCGGACGAGGCGCAATTCGAAATTCGCTT 429

RESULT 3  
 A139650 410 bp mRNA linear EST 08-MAR-2000  
 DEFINITION w942603.x1 NCI CGAP G56 Homo sapiens cDNA clone IMAGE:2473948 3  
 similar to SM:MINK\_HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM  
 CHANNEL PROTEIN ;, mRNA sequence.

ACCESSION A1962650  
 KEYWORDS EST.  
 SOURCE Homo sapiens  
 ORGANISM Human.  
 Biurakova; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.  
 REFERENCE 1 (bases 1 to 410)  
 AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
 Tumor Gene Index  
 JOURNAL Unpublished (1997)  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: c9apbs-remail.nih.gov  
 Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael  
 R. Emmert-Buck, M.D., Ph.D.  
 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima  
 Bonaldi, Ph.D.  
 cDNA Library Arrayed by: Greg Lennon, Ph.D.  
 Arrayed and Sequenced by: National Cancer Sequencing Center  
 Clone Distribution: NCI-CGAP clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LINK at:  
 www.bio.11nl.gov/dbfp/image/image.html  
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 Seq Primer: -400P from 5'bio.

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 /clone IMAGE:2473948  
 /clone\_1ID="NCI CGAP G56"  
 /tissue\_type="pooled germ cell tumors"  
 /lab\_host="DH10B"  
 /note="Vector: p773D-pac (Pharmacia) with a modified  
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 and ss circles were made in vitro. Following HAP purification,  
 this DNA was used as tracer in a subtractive hybridization  
 reaction. The driver was PCR-amplified cDNAs from a pool  
 of 5,000 clones made from the same library (clonesIDB  
 1257096-1258631, 1469064-1470983, and 1475592-1476743).  
 Subtraction by Benito Soares and M. Fatima Bonaldi."  
 BASE COUNT 120 a 35 c 104 t

Query Match 53.7%; Score 393; DB 9; Length 410;  
 Best Local Similarity 100.0%; Pred. No. 1.9e-100;  
 Matches 393; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 110 GACGCTCCGACGACGATTTATCTATATGACAAATGGCGCCGACACCAACACT 169  
 DB 18 GACGCTTCGACGAGATTTATCTATATGACAAATGGCGCCGACACCAACACT 77

OY 170 GACCAAGAGGCGCCGACGACGAAAGTTGACGAGAAATTCATGATGCGATCCGTCG 229  
 DB 78 GACCAAGAGGCGCCGACGACGAAAGTTGACGAGAAATTCATGATGCGATCCGTCG 137

OY 230 CTCATGCTGATGATTTGGAATGTTCTCTTCATCATCGTGGGACATCTGTGAGCACTG 289  
 DB 138 CTCATGCTGATGATTTGGAATGTTCTCTTCATCATCGTGGGACATCTGTGAGCACTG 197  
 OY 290 AATTCAGAAATGCGGACACCTCCATCATCCCTACCCACACGATACATTTGTAGAGACTG 349  
 DB 198 AATTCAGAAATGCGGACACCTCCATCATCCCTACCCACACGATACATTTGTAGAGACTG 257  
 OY 350 CAGGAAAGATGACAGGCAATCTTGAAATCTGAGAGAGGCGCACATCTCATG 409  
 DB 258 CAGGAAAGATGACAGGCAATCTTGAAATCTGAGAGAGGCGCACATCTCATG 317  
 OY 410 AACATGTCGCGCTGGCTTCAAAATGTCCTCCCTGATAGAGGAAAGGACCAACACTTA 469  
 DB 318 AACATTTGGGCGCGCTGGCTTCAAAATGTCCTCCCTGATAGAGGAAAGGACCAACACTTA 377

RESULT 4  
 A1339609 391 bp mRNA linear EST 29-DEC-1998  
 DEFINITION g942607.x1 Soares.NIH/MP.S1 Homo sapiens cDNA clone IMAGE:1935156  
 3' similar to SM:MINK\_HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM  
 CHANNEL PROTEIN ;, mRNA sequence.

ACCESSION A1339609  
 KEYWORDS EST.  
 SOURCE Homo sapiens  
 ORGANISM human.  
 Biurakova; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.  
 REFERENCE 1 (bases 1 to 391)  
 AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
 Tumor Gene Index  
 JOURNAL Unpublished (1997)  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: c9apbs-remail.nih.gov  
 This clone is available royalty-free through LINK ; contact the  
 NCI-CGAP Consortium (http://image.11nl.gov) for further information.  
 Seq Primer: -400P from 5'bio.  
 High quality sequence stop: 380.

FEATURES  
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 /db\_xref="taxon:9606"  
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 /clone\_1ID="Soares.NIH/MP.S1"  
 /tissue\_type="Human melanocyte, fetal heart, and  
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 /lab\_host="DH10B"  
 /note="Organ: mixed (see below); Vector: p773D-pac  
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 normalized libraries (melanocyte 2MBB, pregnant uterus  
 NBH90 and fetal heart NBH159) were added, and 88 circles  
 were made in vitro. The driver was PCR-amplified cDNAs  
 from a pool of 5,000 clones made from the same 3 libraries. The pools  
 consisted of I.M.A.G.E. clones 260232-265223,  
 340488-345479, and 484488-489479."  
 BASE COUNT 119 a 93 c 86 t

Query Match 52.7%; Score 385.8; DB 9; Length 391;  
 Best Local Similarity 99.5%; Pred. No. 2.1e-98;  
 Matches 387; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 111 ACGCTTCGAGAGATTTATCTATATGACAAATGGCGCCGACACCAACCAACTG 170

Db 3 ACAGCTTCGAGAGATTTTATCTATGTTGGGACATTGGGCGGACAAACACAGACCTG 62  
 OY 171 AGCAAGAGGCGCTCCAGACGCAAACTGTGCTGGAGAACTCTTCAATGATCTCCGTGAC 230  
 Db 63 AGCAAGAGGCGCTCCAGACGCAAACTGTGCTGGAGAACTCTTCAATGATCTCCGTGAC 122  
 OY 231 TCATGTGTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 290  
 Db 123 TCATGTGTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 182  
 OY 291 AATCCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 350  
 Db 183 AATCCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 242  
 OY 351 AGCAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 410  
 Db 243 AGCAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 302  
 OY 411 ACATGTGTGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGG 470  
 Db 303 ACATGTGTGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCT 362  
 OY 471 ATGCGACGTCCAGACATGAGAGATGCCA 499  
 Db 363 ATGCGACGTCCAGACATGAGAGATGCCA 391

RESULT 5  
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 DEFINITION A1246239.1 Searles JHMPV, USA Homo sapiens CDNA clone IMAGE:1857942  
 CHANNEL PROTEIN ; mRNA sequence.  
 ACCESSION A1246239  
 VERSION A1246239.1 GI:3841636  
 KEYWORDS EST.  
 SOURCE human.  
 ORGANISM Homo sapiens; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.  
 REFERENCE 1 (bases 1 to 372)  
 NC-CAP http://www.ncbi.nlm.nih.gov/ncicgap.  
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
 Tumor Gene Index  
 JOURNAL Unpublished (1997)  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 National Cancer Institute  
 This clone is available royalty-free through NCI. Contact the  
 IMAGE Consortium (info@image.llnl.gov) for further information.  
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 High quality sequence stop: 365.  
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 pregnant uterus"  
 /lab\_protocol="DH108"  
 /vector="pUC19" (see below): Vector: pUT7AD-pc  
 (Pharmacia) with a modified polylinker site. NheI  
 site 2: Eco RI; BstI amount of plasmid DNA from three  
 normalized libraries (melanocyte 2NBHM, pregnant uterus  
 JHMPV, and fetal heart JHMH19W) were mixed, and ss circles  
 were made in vitro. Following HAP purification, this DNA  
 was used as template in a subtractive hybridization  
 reaction. The driver was PCR-amplified cDNAs from pools of  
 500 cells. The subtracted cDNA was ligated into a  
 construct of pMA-CAT, clone#260232-265232, The pools  
 340488-345479, and 484488-489479."

BASE COUNT 115 a 89 c 86 g 82 t  
 ORIGIN  
 Query Match 50.8%; Score 372; PB 9; Length 372;  
 Best local similarity 100.0%; 127e-94;  
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 OY 178 GCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 237  
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 OY 238 GATGATGAGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 297  
 Db 121 GATGATGAGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 180  
 OY 298 GAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 357  
 Db 181 GAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 240  
 OY 358 GTACAGAGCAATCTGATCTATGAGAAATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 417  
 Db 241 GTACAGAGCAATCTGATCTATGAGAAATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 300  
 OY 418 TCGCGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCT 477  
 Db 301 TCGCGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCT 360  
 OY 478 GTCCAGACATGGA 489  
 Db 361 GTCCAGACATGGA 372  
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 DEFINITION A2693989.1 Searles JHMPV, USA Homo sapiens genomic 5', DNA sequence.  
 ACCESSION A2693989  
 VERSION A2693989.1 GI:11878908  
 KEYWORDS GSS.  
 SOURCE human.  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.  
 REFERENCE 1 (bases 1 to 350)  
 NC-CAP http://www.ncbi.nlm.nih.gov/ncicgap.  
 TITLE Exon-trap tags from a HL-60 Genomescreen(TM) Library  
 JOURNAL Unpublished (2000)  
 COMMENT Contact: Greg Henkel  
 Gene Expression  
 National Cancer Institute  
 This clone is available royalty-free through NCI. Contact the  
 IMAGE Consortium (info@image.llnl.gov) for further information.  
 Insert Length: 921 Std Error: 0.00  
 Seq primer: -400D from Gdbco  
 High quality sequence stop: 365.  
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 /db\_xref="taxon:9606"  
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 pregnant uterus"  
 /lab\_protocol="DH108"  
 /vector="pUC19" (see below): Vector: pUT7AD-pc  
 (Pharmacia) with a modified polylinker site. NheI  
 site 2: Eco RI; BstI amount of plasmid DNA from three  
 normalized libraries (melanocyte 2NBHM, pregnant uterus  
 JHMPV, and fetal heart JHMH19W) were mixed, and ss circles  
 were made in vitro. Following HAP purification, this DNA  
 was used as template in a subtractive hybridization  
 reaction. The driver was PCR-amplified cDNAs from pools of  
 500 cells. The subtracted cDNA was ligated into a  
 construct of pMA-CAT, clone#260232-265232, The pools  
 340488-345479, and 484488-489479."







Db	325	AGAGATCGATGTCGATGCGAGATCGACATCGGATCATGTCATGTAACCTGGGGGCG	384
Qy	422	CGTGGGTTCACAAATGTCCTCCCTGATTAAGCA	452
Db	385	ACGGGATTCACGATGACACCGCTGATTAAGCA	415
RESULT 9			
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DEFINITION	Mus musculus cDNA clone IMAGE:4481325 5'		linear
ACCESSION	BC626195		
VERSION	BC626195.1		
KEYWORDS	EST.		
ORIGIN			
SOURCE	house mouse.		
ORGANISM	Mus musculus		
REFERENCE	Elkayrola, Metazon; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.		
AUTHORS	1. NIH-MGC http://mgs.scrib.nhl.gov/		
TITLE	National Institutes of Health, Mammalian Gene Collection (MGC)		
JOURNAL	Unpublished (1999)		
COMMENT	Contact: Robert Strauberg, Ph.D. Email: cgaops-remail.nhl.gov Tissue Procurement: The Cepko Laboratory cDNA Library Preparation: Life Technologies, Inc. Library Stripped by: The I.M.A.G.E. Consortium (LLMI) Library Sequenced by: The I.M.A.G.E. Consortium DNA Sequencing: The I.M.A.G.E. Consortium Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLMI at: http://image.llnl.gov Plate: LLMI0316 row 5 column 22 High quality sequence stop: 535.		
FEATURES			
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	/tissue_type="retina"		
	/lab_host="DH10d (phage-resistant)"		
	/lib_name="Ogan: eye; vector: pCMV-SK(+)6; size: 3. kb; Strategy: PCR; insert: 3 kb; Library: full-length clones and constructed by Life Technologies. Note: this is a NIH-MGC Library."		
BASE COUNT	249 a 218 c 302 g 234 t		
ORIGIN			
Query Match	35 1% Score 256.6; DB 12; Length 1003;		
Best Local Similarity	80.8% Overlap 67.6%		
Matches 324; Conservative	0; Mismatches 74; Indels 3; Gaps 2;		
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Db	78	GCAAGGAGGAGACATGCGCACATTTAGCACACGAGACATGAGAGATGCTCTTCA	137
Qy	121	AAGCATTTTATCTACTATATGAGACATGTGGCGCCACACACANCGTACGACAGGCG	180
Db	138	AAAGATTTTATCTATATATGAGACATGTGGCGCCACACACACGCGACAGAGAGCG	197
Qy	181	CTTCCACGACCAATGATATCTGAGAAATCTGTACATGTCATCTGTTACCTGATGAT	240
Db	198	ACTCCAGCGCAAGATGATGACGAGAACTCTCAAGACATCTGTAACCTGATAGTAT	257
Qy	241	GATGGAATGCTCTTCTATCATCATGTCGACATCTCTGATGACATGTGAATCTAAG	300
Db	258	GGGACATGCTCTTCTATCATCATGTCGACATCTCTGATGACATGTGAATCTAAG	317
Qy	301	ACGGAGACATCGACACCGCTACACACATCTCTATGACATGAGATGACAGAAATGA	360
Db	318	CGGAGACATCGACACCGCTACACACATCTCTATGACATGAGATGACAGAAATGA	377

Oy	361	CAGACCCAAATCTGATTTCATGTGGAGAATGCGA- -GCCCAACTCCTCATCGAACAAT-TGC 417 
Dy	378	GAAAATGTACATTTCTCATCTTGAAAAAGTCCCAGAACGCCACCACCAATCATCATGAGAAATGGGG 437 
Oy	418	TGCGGCTGTGGTTTTCAAATATCCCCCGTAGTAAGAGCACAAAGG 458 
Dy	438	GCGACAGGGGTTTCACAGTGTCCACCCCTTATTTGATAAAGAAATG 478 
RESULT 10		
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DEFINITION	Atherosys RAGE Library Homo sapiens cDNA, mRNA sequence.	
DESCRIPTION		
ACCESSION	BG221966	JGI:J3747987
VERSION	BG221966.1	
KEYWORDS	EST.	
SOURCE	human.	
ORGANISM	Homo sapiens	
REFERENCE	Rodriguez-Medacozz; Chorclata; Crutclata; Vertebrate; Euteleostomi; Rodriguez-Medacozz; Primates; Catarrhini; Hominoidea; Homo. 1. (bases 1 to 746) Harrington J.J., Sherif B., Rundlett S., Jackson P.D., Perry R., Calis S., Leventhal C.C., Thornton M., Ramchandran K., Whittington J., Lerner L.G., Costanzo D., McGillivray K., Booser S., Mayes R., Smith E., Veloso N., Kilka A.R., Hess J., Colbran K., Lo K., Offenbacher J.E., Danzig Y. and Duzy G.M. Creation of genome-wide protein expression libraries using random Nbt. Biotechnol. 19 (5), 440-445 (2001)	
JOURNAL MEDLINE COMMENT	Contact: Scott J. Cain 3201 Carnegie Ave, Cleveland, OH 44115, USA Tel.: 216 431 9900 Fax: 216 431 9586 Email: scott.j.cain@atherysys.com	
FEATURES		
source	Location/Qualifiers  1..746 -/organism= "Homo sapiens" -/db_xref=taxon:9606" -/gene_lib=Athersys RAGE Library"	
Description: Cell-line: The production of Genome-Wide Protein Expression Libraries using Random Activation of Gene Expression". Nature Biotechnology, In press. Note that even though the cell type indicated is HT1080, since a random activation method was used, those sequence tags are not necessarily expressed in HT1080 under normal circumstances."		
BASE COUNT      241 a    138 c    123 g    243 t     1 others		
ORIGIN		
Query Match                  34.3% ; Score 250.8 ; DB: 12; Length 746;		
Best Local Similarity        Pred. No. 5,6e+00 ;		
Matches 280; Conservative    0; Mismatches 13; Indels    2; Gaps    2;		

QY	241	GATTGGAAATGTCCTTCCTTCATG-CATGCTGGGCGCATCCGCTGAGCATGTGTAATC	294
Db	692	GATTGGAAATGTCCTTCCTTCATGAAATGAGGGGCGCATCTGGGAGCACTGTGAATC	746
RESULT 11			
LOCUS			
DEFINITION			
ACCESSION			
KEYWORDS			
SOURCE			
ORGANISM			
REFERENCE			
AUTHORS			
TITLE			
JOURNAL			
COMMENT			
FEATURES			
SOURCE			

Db	121	GATGCGAGGGCCACGAGGCAGAAATTCATAATGTCTTCTTGTAAGAAGAGGATGCTCTT	180
Oy	553	CTCTTTGTGTGAAGATTTCATCATGAGATATGTGCTGTGGCAATTAAGATNAGTACATT	612
Db	191	CCTCCTGTGTGAAGATTTCATCATGAGATATGTGCTGTGGCAATTAAGATNAGTACATT	240
Oy	613	CAM	615
Dy	241	CAM	243
RESULT 12			
LOCUS	404		
DEFINITION			
ACCESSION	AA633404	188 bp	mRNA EST 28-OCT-1997
VERSION	np65911.61	NCI CGAP Bt2	Homo sapiens cDNA clone IMAGE:1131621.3'
KEYWORDS			similar to SW-MYC_HUMAN P15382; ISF SLOW VOLUME-GATED POTASSIUM CHANNEL PROTEIN ; ; mRNA sequence.
SOURCE	AA633404		
ORGANISM	AA633404.1	GI:2555264	
EST			
Protein			
Homologs			
Eukaryotic:	Metazoa:	Chordata:	Crinata:
			Vertebrata:
			Euteleostomi:
	Mammalia:	Eutheria:	Primates:
			Catarrhini:
			Hominoidea:
			Homo-
			(Bases 1 to 188)
REFERENCE	NCI-CGAP	http://www.ncbi.nlm.nih.gov/ncicgap	
AUTHORS	National Cancer Institute,	Cancer Genome Anatomy Project (CGAP),	
TITLE	Human Gene Mapped to Chromosome 12 (1997).		
JOURNAL	Contact: Robert Strimberg,	Ph.D.	
COMMENT	Email: cgaps@femail.nih.gov		
	Tissue Procurement: Christopher Moskalko,	M.D., Ph.D., Michael R.	
	Emmert-Buck,	M.D., Ph.D.	
	cDNA Library Preparation: M. Bento Soares,	Ph.D.	
	Library Arrayed by: Greg Lennon,	Ph.D.	
	RNA Sequencing: David Winkler, CGP	Genome Sequencing Center	
	Chromosomal Localization Information can be found through the I.M.A.G.E. Consortium/LDN at:		
	www.bio.llnl.gov/dbcp/image.html		
	Insert Length: 785	Std Error: 0.00	
	Seq Primer: -40m13 fwd. ET from Amsterdam		
	High quality sequence #top: 167.		
FEATURES			
source			
	Location/Qualifiers		
	1..organelle=Homo sapiens*		
	/db_xref=taxon:9606*		
	/clone=IMAGE:1131621*		
	/clone.lib=NCI CGAP Bt2*		
	/sex=female, pooled*		
	/tissue_type=breast*		
	/lab.name=DRI db/r73D-PKC (pharmacla) with a modified		
	polylinker. The strand cDNA was prepared from pooled bulk		
	breast tumor tissue, and was then primed with a Not I -		
	oligo(dT) primer. Double-stranded cDNA was ligated to Eco		
	RI adaptors (pharmacla), digested with Not I and cloned		
	into the Not I and Eco RI sites of the modified pRTT3		
	vector. This library is the normalized version of		
	NCI CGAP Bt1.1 library was constructed by Bento Soares		
	and M. Estrela.		
	66 a	45 e	32 t
BASE COUNT			
ORIGIN			
Query Match	25.7%	Score 188;	DB 9; Length 188;
Best Local Similarly	100.0%;	Fwd. No. 1; 9e-42;	
Matches 188;	Conservative	0; Mismatches	0; Indels
		0; Gaps	0;
Oy	286	TGTCAATTCAGACGACGCAATTCATCAATGACCTGTACGCCATGCTGTACAGCA	345
Dy	1	TGTCAATTCAGACGACGCAATTCATCAATGACCTGTACGCCATGCTGTACAGCA	60
346	CTGTGACGAGAAAAGACAAGCACCAATTCGATCTGTGAAATATCGAAGGCACACTATCA	405	

Db 61 CTGGCAGGAAAGATACAGAGCCAAATCTGATAGTAAGATCGAAGCCACCATCA 120  
 QY 406 TGAAGACATGTGTGCGGCTGTGGCTTAAATGTCCCTCCGTATAGGGAGAAAGGCCCAAG 465  
 Db 121 TGAAGACATGTGTGCGGCTGTGGCTTAAATGTCCCTCCGTATAGGGAGAAAGGCCCAAG 180  
 QY 466 CTAAACATC 473  
 Db 181 CTAAACATC 188  
 RESULT 13  
 B638225  
 LOCUS 1Ab014142 Bovine Abomasum cDNA Library Bos taurus cDNA 5', mRNA  
 DEFINITION  
 sequence.  
 B638225 351 bp mRNA linear EST 11-JUN-2001  
 VERSION B638225.1 GI:14337597  
 ACCESSION  
 SOURCE  
 ORGANISM  
 Bos taurus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;  
 Bovidae; Bovinae; Bos.  
 1 (bases 1 to 351)  
 Moore,S.S., Hansen,C., Li,C., Fu,A., Meng,Y. and Li,G.  
 Molecular cloning of a bovine abomasum tissue  
 cDNA library (2001)  
 Unpublished  
 Contact: Dr. Stephen Moore  
 Beef Genomics Laboratory  
 Dept of AFS, University of Alberta  
 410 Agri/For, Dept of AFMS, U of A, Edmonton, AB, T6G 2P5, Canada  
 Tel.: 780 492 0169  
 Fax: 780 492 4285  
 Email: s.moore@ualberta.ca  
 The sequence best matches gb|AF001719 (Homo sapiens genomic DNA,  
 chromosome 21q, section 63/105) in main database at high score of  
 212.0 and E-value of 9e-53  
 PCR Primers  
 FORWARD: M13 Reverse  
 BACKWARD: M13 Forward  
 Seq primer: 13 Primer  
 High quality sequence stop: 351  
 POLY-A-NO  
 Location/Qualifiers  
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 /db\_xref="taxon:9913"  
 /clone\_1lb="Bovine Abomasum cDNA Library"  
 /note="male and one female mixed  
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 /cell\_type="Epithelial"  
 /dev\_stage="Young adult"  
 /lab\_host="XL1-blueMR"-strain"  
 /note="Organ: Abomasum; Vector: Uni-ZAPXR; Site:1; EcoR  
 I; Site:2; Xho I"  
 BASE COUNT 104 a 89 c 76 g 82 t  
 ORIGIN  
 Query Match 24.0%; Score 175.4; DB 13; Length 351;  
 Best Local Similarity 86.5%; Pred.No.9.5e-39;  
 Matches 205; Conservative 0; Mismatches 31; Indels 1; Gaps 1;

Db 235 CTGCAAGCCAGGTTGATGATGAGATCTTCTACTATGATCTTATGATGATG 294  
 QY 242 ATTGGAATGTCTCTTT CATGATCTGTGGCATCCCTGTGGAGACATGTGAATTCAA 297  
 Db 295 ATCGGAATGTCTCTTCTCAATGATTTGACCATCTGTGTGAGCAGCTGAATTCAA 351  
 RESULT 14  
 AA935321  
 LOCUS AA935321 121 bp mRNA linear EST 07-JUL-1998  
 DEFINITION  
 o071909.s1 NCI-CGAP G24 Homo sapiens cDNA clone IMAGE:1571680 3'  
 similar to SK-MIN6 HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM  
 CHANNEL PROTEIN ; mRNA sequence.  
 AA935321  
 VERSION AA935321.1 GI:3092478  
 KEYWORDS  
 EST.  
 SOURCE  
 ORGANISM  
 human.  
 Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homini; Homo.  
 REFERENCE  
 1 NCI-CGAP http://www.ncbi.nlm.nih.gov/ncigap.  
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
 Tumor Gene Index  
 Unpublished (1997)  
 CONTACT: Robert Strauszberg, Ph.D.  
 Email: cgaap-r@mail.nih.gov  
 Tissue Procurement: Christopher A. Koskialuk, M.D., Ph.D., Michael  
 Biesecker  
 cDNA Library Preparation: M. Bento Soares, Ph.D.  
 cDNA Library Arrayed by: Greg Lennon, Ph.D.  
 DNA Sequencing by: Washington University Genome Sequencing Center  
 Clone distribution: NCI-CGAP clone distribution information can be  
 found through the I.M.A.G.E. Consortium/ILM, at:  
 www.bio.11nl.gov/dbfp/image/image.html  
 Trace considered overall poor quality  
 Insert length: 763 Std Error: 0.00  
 Seq primer: -40ml3 fwd. ET from Amersham  
 High quality sequence stop: 1.  
 Location/Qualifiers  
 1. 121  
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 /db\_xref="taxon:9606"  
 /clone\_1lb="NCI-CGAP G24"  
 /tissue\_type="Pooled germ cell tumors"  
 /lab\_host="DH10B"  
 /note="Vector: pTV3D-Pac (Pharmacia) with a modified  
 polylinker. 1st strand cDNA was prepared from 3 pooled  
 germ cell tumors, and was then primed with a Not I to Eco  
 RI adaptor (Pharmacia) and digested with Not I and cloned  
 into the Not I and Eco RI sites of the modified pTV3D  
 vector. Library is normalized. Library was constructed by  
 Bento Soares and M. Felima Bonaldo."  
 BASE COUNT 33 a 31 c 28 g 29 t  
 ORIGIN  
 Query Match 16.3%; Score 119.4; DB 9; Length 121;  
 Best Local Similarity 99.2%; Pred.No.4.4e-23;  
 Matches 120; Conservative 0; Mismatches 1; Indels 0;



GenCore version 5.1.4.p5.4578  
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OM protein - protein search, using sw model

Run on: May 15, 2003, 14:23:09 ; Search time 14 Seconds

(without alignments)

258,501 Million cell updates/sec

Title: US-09-550-163-2

Perfect score: 62

Sequence: 1 NGLSLNFRQLIEDVFRRII.....EESKATIHENIGAGKTKSP 123

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 262574 seqs, 29422922 residues

Total number of hits satisfying chosen parameters: 262574

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing filter 43 summaries

Database : Issued.Patents.AA:\*

1: /sgn2.6/pdata/1/1aa/5A.COMB.pep:\*

2: /sgn2.6/pdata/1/1aa/5B.COMB.pep:\*

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4: /sgn2.6/pdata/1/1aa/5B.COMB.pep:\*

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6: /sgn2.6/pdata/1/1aa/5B.COMB.pep:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Match	Query length	DB ID	Description
1	132.5	21.0	129	3	US-09-069-896-3
2	132.5	21.0	129	4	US-09-135-021-78
3	132.5	21.0	129	4	US-09-115-020-4
4	132.5	21.0	129	4	US-09-115-020-4
5	132.5	21.0	129	4	US-09-115-020-4
6	132.5	21.0	129	4	US-09-115-020-4
7	132.5	21.0	129	4	US-09-115-020-4
8	132.5	21.0	129	4	US-09-115-020-4
9	132.5	21.0	129	4	US-09-115-020-4
10	132.5	21.0	129	4	US-09-115-020-4
11	132.5	21.0	129	4	US-09-115-020-4
12	132.5	21.0	129	4	US-09-115-020-4
13	132.5	21.0	129	4	US-09-115-020-4
14	132.5	21.0	129	4	US-09-115-020-4
15	132.5	21.0	129	4	US-09-115-020-4
16	132.5	21.0	129	4	US-09-115-020-4
17	132.5	21.0	129	4	US-09-115-020-4
18	132.5	21.0	129	4	US-09-115-020-4
19	132.5	21.0	129	4	US-09-115-020-4
20	132.5	21.0	129	4	US-09-115-020-4
21	132.5	21.0	129	4	US-09-115-020-4
22	132.5	21.0	129	4	US-09-115-020-4
23	132.5	21.0	129	4	US-09-115-020-4
24	132.5	21.0	129	4	US-09-115-020-4
25	132.5	21.0	129	4	US-09-115-020-4
26	132.5	21.0	129	4	US-09-115-020-4
27	132.5	21.0	129	4	US-09-115-020-4

28	65	10.3	1835	4	US-09-404-650-5
29	65	10.3	2175	4	US-09-404-650-5
30	65	10.3	2188	4	US-09-404-650-5
31	64.5	10.2	970	2	US-08-673-789-7
32	64.5	10.2	972	1	US-08-102-809-10
33	64.5	10.2	975	1	US-08-102-809-10
34	64.5	10.2	976	4	US-09-511-477-4
35	64.5	10.2	976	4	US-09-511-507-4
36	64.5	10.2	1367	2	US-08-249-687C-2
37	64	10.1	1367	2	US-08-625-819-2
38	64	10.1	1367	3	US-08-746-559A-2
39	63.5	10.1	1367	4	US-08-964-641B-18
40	63.5	10.0	439	3	US-08-448-722A-5
41	63.5	10.0	439	3	US-08-448-722A-5
42	63.5	10.0	439	4	US-08-189-309B-5
43	63.5	10.0	2332	1	US-08-276-594A-2
44	63.5	10.0	2351	1	US-08-366-851A-2
45	63.5	10.0	2351	6	517184A-2

## ALIGNMENTS

RESULT 1  
US-09-069-896-3  
Sequence 3, Application US/09069896  
Patent No. 6071720  
GENERAL INFORMATION:  
APPLICANT: Genentech, Inc., Jennifer L.  
APPLICANT: Patterson, Chandra  
TITLE OF INVENTION: DELAYED RECTIFIER POTASSIUM  
NUMBER OF SEQUENCES: 4  
CORRESPONDENCE ADDRESS:  
STREET: 1174 Center Drive  
CITY: Palo Alto  
STATE: CA  
COUNTRY: USA  
ZIP: 94304  
COMPUTER READABLE FORM:  
SEQUENCE TYPE: Nucleic Acid  
COMPUTER SYSTEM COMPATIBLE  
OPERATING SYSTEM: DOS  
SOFTWARE: FASTSEQ for Windows Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/069, 896  
FILING DATE:  
PUBLICATION DATA:  
CLASSIFICATION:  
PRIORITY NUMBER:  
FILING DATE:  
ATTORNEY/AGENT INFORMATION:  
NAME: Carrone, Michael C  
REGISTRATION NUMBER: 39,132  
REFERENCE/DOCKET NUMBER: PF-0507 US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 650-945-5205  
TELEFAX: 650-945-4166  
TELEX:  
INFORMATION FOR SEQ ID NO: 3:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 129 amino acids  
TYPE: amino acid  
STRANDEDNESS: single  
ORIENTATION: 5' to 3'  
IMMEDIATE SOURCE:  
LIBRARY: Genbank  
CLONE: 452497  
US-09-069-896-3

Query Match

21.0%; Score 132.5; DB 3; Length 129;

Best Local Similarity 45.18; Pred. No. 9.3e-08;  
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;  
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DB 45 LTVLWVWGFEFTLGLMSTYINRSKRLHNSNDPNVYTESDAMQKDAVY 95

RESULT 2  
US-09-135-021-78  
Sequence 78; Application US/09135021A  
Patent No. 6150104  
GENERAL INFORMATION:  
APPLICANT: Splawski, Igor  
APPLICANT: Keating, Mark T.  
TITLE OF INVENTION: CAUSE ARRYHYTHIA SUSCEPTIBILITY IN KVLOT1 WHICH CAUSES JERVELL  
FILE REFERENCE: 2323-128  
CURRENT APPLICATION NUMBER: US/09/135.021A  
EARLIER FILING DATE: 1998-08-17  
EARLIER FILING DATE: 1997-06-13/094,655  
EARLIER FILING DATE: 1997-06-13/094,477  
EARLIER FILING DATE: 1998-07-29  
NUMBER OF SEQ ID NOS: 80  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 78  
LENGTH: 129  
TYPE: CDS  
ORGANISM: Homo sapiens  
US-09-135-021-78

Query Match 21.0%; Score 132.5; DB 4; Length 129;  
Best Local Similarity 45.18; Pred. No. 9.3e-08;  
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;  
OY 51 LTVLWVWGFEFTLGLMSTYINRSKRLHNSNDPNVYTESDAMQKDAVY 100  
||||| : : : : : ||| : : : : : ||| : : : : :  
DB 45 LTVLWVWGFEFTLGLMSTYINRSKRLHNSNDPNVYTESDAMQKDAVY 95

RESULT 3  
US-09-135-020-4  
Sequence 4; Application US/09135020  
Patent No. 6132026  
GENERAL INFORMATION:  
APPLICANT: Keating, Mark T.  
APPLICANT: Sanguinetti, Michael C.  
TITLE OF INVENTION: MUTATIONS IN THE KCNE1 GENE ENCODING HUMAN MLK WHICH  
TITLE OF INVENTION: CAUSE ARRYHYTHIA SUSCEPTIBILITY THEREBY ESTABLISHING  
FILE REFERENCE: 2323-131  
CURRENT APPLICATION NUMBER: US/09/135.020  
EARLIER FILING DATE: 1998-08-17  
EARLIER FILING DATE: 1997-08-29  
EARLIER FILING DATE: 1996-10-29  
EARLIER FILING DATE: 1995-12-22  
EARLIER FILING DATE: 1995-12-22  
EARLIER FILING DATE: 1998-07-29  
NUMBER OF SEQ ID NOS: 114  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 4  
LENGTH: 129  
TYPE: CDS  
ORGANISM: Homo sapiens  
US-09-135-020-4

Query Match 21.0%; Score 132.5; DB 4; Length 129;  
Best Local Similarity 45.18; Pred. No. 9.3e-08;

Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;  
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DB 45 LTVLWVWGFEFTLGLMSTYINRSKRLHNSNDPNVYTESDAMQKDAVY 95

RESULT 4  
US-09-135-010A-4  
Sequence 4; Application US/09135010A  
Patent No. 6132026  
GENERAL INFORMATION:  
APPLICANT: Keating, Mark T.  
APPLICANT: Sanguinetti, Michael C.  
APPLICANT: Curran, Mark E.  
APPLICANT: Landes, Gregory M.  
APPLICANT: Comoros, Timothy D.  
APPLICANT: Sanguinetti, C.  
TITLE OF INVENTION: KVLOT1 - A LONG QT SYNDROME GENE  
FILE REFERENCE: 2323-133  
CURRENT APPLICATION NUMBER: US/09/135.010A  
EARLIER FILING DATE: 1998-08-17  
EARLIER FILING DATE: 1997-08-29  
EARLIER FILING DATE: 1997-08-29  
EARLIER FILING DATE: 1997-08-29  
EARLIER FILING DATE: 1996-10-29  
EARLIER FILING DATE: 1995-12-22  
EARLIER FILING DATE: 1995-12-22  
NUMBER OF SEQ ID NOS: 115  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 4  
LENGTH: 129  
TYPE: CDS  
ORGANISM: Homo sapiens  
US-09-135-010A-4

Query Match 21.0%; Score 132.5; DB 4; Length 129;  
Best Local Similarity 45.18; Pred. No. 9.3e-08;  
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;  
OY 51 LTVLWVWGFEFTLGLMSTYINRSKRLHNSNDPNVYTESDAMQKDAVY 100  
||||| : : : : : ||| : : : : : ||| : : : : :  
DB 45 LTVLWVWGFEFTLGLMSTYINRSKRLHNSNDPNVYTESDAMQKDAVY 95

RESULT 5  
US-09-444-871-4  
Sequence 4; Application US/09444871  
Patent No. 6132026  
GENERAL INFORMATION:  
APPLICANT: Keating, Mark T.  
APPLICANT: Sanguinetti, Michael C.  
TITLE OF INVENTION: MUTATIONS IN THE KCNE1 GENE ENCODING HUMAN MLK WHICH  
TITLE OF INVENTION: CAUSE ARRYHYTHIA SUSCEPTIBILITY THEREBY ESTABLISHING  
FILE REFERENCE: 2323-131  
CURRENT APPLICATION NUMBER: US/09/444.871  
EARLIER FILING DATE: 1998-11-22  
EARLIER FILING DATE: 1998-08-17  
EARLIER FILING DATE: 1998-08-17  
EARLIER FILING DATE: 1997-08-29  
EARLIER FILING DATE: 1996-10-29  
EARLIER FILING DATE: 1995-12-22  
EARLIER FILING DATE: 1995-12-22  
EARLIER FILING DATE: 1998-07-29  
NUMBER OF SEQ ID NOS: 114  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 4  
LENGTH: 129  
TYPE: CDS  
ORGANISM: Homo sapiens  
US-09-444-871-4



SOFTWARE: PatentIn Ver. 2.0  
 SEQ ID NO 4  
 LENGTH: 129  
 ORGANISM: Homo sapiens  
 US-09-444-295.4

Query Match 21.0%; Score 132.5; DB 4; Length 129;  
 Best Local Similarity 45.1%; Pred. No. 9.3e-08;  
 Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

51 LTVLWVGPFPGFTLGLMISTVSKRSRRENSDPHYIVED-  
 LTVLWVGPFPGFTLGLMISTVSKRSRRENSDPHYIVESDAMQERKAVY 95

US-09-597-735-4

Sequence 4, Application US/09597735  
 Patent No. 6432687  
 GENERAL INFORMATION:  
 APPLICANT: Keating, Mark T.  
 APPLICANT: Sanguinetti, Michael C.  
 APPLICANT: Curran, Mark E.  
 APPLICANT: Landes, Gregory M.  
 APPLICANT: Connors, Timothy D.  
 APPLICANT: Barn, Timothy C.  
 TITLE OF INVENTION: KNOT1 - A LONG QT SYNDROME GENE  
 FILE REFERENCE: 2323-133  
 CURRENT APPLICATION NUMBER: US/09/597,735

EARLIER FILING DATE: 2000-06-19  
 EARLIER APPLICATION NUMBER: 09/135,010  
 EARLIER FILING DATE: 1998-08-29  
 EARLIER APPLICATION NUMBER: 08/994,477  
 EARLIER FILING DATE: 1998-07-29  
 EARLIER APPLICATION NUMBER: 08/922,068  
 EARLIER FILING DATE: 1997-08-29  
 EARLIER APPLICATION NUMBER: 08/739,383  
 EARLIER FILING DATE: 1996-10-29  
 EARLIER APPLICATION NUMBER: 60/019,014  
 NUMBER OF SEQ ID NOS: 11  
 SOFTWARE: PatentIn Ver. 2.0  
 SEQ ID NO 4  
 LENGTH: 129  
 TYPE: PRT  
 ORGANISM: Homo sapiens  
 US-09-597-735-4

Query Match 21.0%; Score 132.5; DB 4; Length 129;  
 Best Local Similarity 45.1%; Pred. No. 9.3e-08;  
 Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

51 LTVLWVGPFPGFTLGLMISTVSKRSRRENSDPHYIVED-  
 LTVLWVGPFPGFTLGLMISTVSKRSRRENSDPHYIVESDAMQERKAVY 95

RESULT 7

US-09-444-295-4  
 Sequence 4, Application US/09444295  
 Patent No. 6432644  
 GENERAL INFORMATION:  
 APPLICANT: Keating, Mark T.  
 APPLICANT: Sanguinetti, Michael C.  
 APPLICANT: Curran, Mark E.  
 APPLICANT: Landes, Gregory M.  
 APPLICANT: Connors, Timothy D.  
 APPLICANT: Barn, Timothy C.  
 TITLE OF INVENTION: KNOT1 - A LONG QT SYNDROME GENE  
 FILE REFERENCE: 2323-133  
 CURRENT APPLICATION NUMBER: US/09/444,295  
 CURRENT FILING DATE: 1999-11-22

PRIOR APPLICATION NUMBER: 09/135,020  
 PRIOR FILING DATE: 1998-08-17  
 PRIOR APPLICATION NUMBER: 08/921,068  
 PRIOR FILING DATE: 1997-08-29  
 PRIOR APPLICATION NUMBER: 08/739,383  
 PRIOR FILING DATE: 1998-07-29  
 PRIOR APPLICATION NUMBER: 60/019,014  
 PRIOR FILING DATE: 1995-12-22  
 PRIOR APPLICATION NUMBER: 60/094,477  
 PRIOR FILING DATE: 1998-07-29  
 NUMBER OF SEQ ID NOS: 114  
 SOFTWARE: PatentIn Ver. 2.0  
 SEQ ID NO 4  
 LENGTH: 129  
 TYPE: PRT  
 ORGANISM: Homo sapiens  
 US-09-444-295-4

Query Match 21.0%; Score 132.5; DB 4; Length 129;  
 Best Local Similarity 45.1%; Pred. No. 9.3e-08;  
 Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

51 LTVLWVGPFPGFTLGLMISTVSKRSRRENSDPHYIVED-  
 LTVLWVGPFPGFTLGLMISTVSKRSRRENSDPHYIVESDAMQERKAVY 95

RESULT 8

US-09-471-468-3  
 Sequence 3, Application US/09471468  
 Patent No. 6432687  
 GENERAL INFORMATION:  
 APPLICANT: Hillman, Jennifer L.  
 APPLICANT: Patterson, Chandra  
 APPLICANT: Portley, Neil  
 APPLICANT: Portley, Neil  
 TITLE OF INVENTION: CHANNEL HOMOLOC  
 NUMBER OF SEQUENCES: 4  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Incyte Pharmaceuticals, Inc.  
 STREET: 3174 Porter Drive  
 CITY: Palo Alto  
 STATE: CA  
 COUNTRY: USA  
 ZIP: 94304  
 COMPUTER READABLE FORM:  
 MEDIUM TYPE: Diskette  
 COMPUTER: IBM Compatible  
 OPERATING SYSTEM: DOS  
 SOFTWARE: FASTSEQ for Windows Version 2.0  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/09/471,468  
 FILING DATE:  
 CLASSIFICATION:  
 PRIOR APPLICATION DATA:  
 APPLICATION NUMBER: 09/069,896  
 FILING DATE:  
 ATTORNEY/AGENT INFORMATION:  
 NAME: [REDACTED]  
 REGISTRATION NUMBER: 39,112  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: 650-855-0555  
 TELEFAX: 650-845-4166  
 TEXT: FOR SEQ ID NO: 3:  
 SOURCE CHARACTERISTICS:  
 LENGTH: 129 amino acids  
 TYPE: amino acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 IMMEDIATE SOURCE:  
 LIBRARY: Genbank

CLONE: 452497  
US-09-471-468-3

Query Match 21.0% Score 132.5; DB 4; Length 129;  
Best Local Similarity 45.1% Pred. No. 9.3e-08;  
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

OY 51 LYLAWMGESFVIALVSTVSKRRRHSNDPYQIYVED-WQERYKSQI 100  
DB 45 LYLAWLGFGEFTLGLIMLSTIRSKRLHSNDPNVYTESDAMQEKQNAV 95

RESULT 9  
US-09-597-732-4

Sequence 4, Application US/09597732

Patent No. 6451334  
INVENTOR: Mark T.  
APPLICANT: Sanguinetti, Michael C.  
APPLICANT: Curran, Mark E.  
APPLICANT: Landes, Gregory M.  
APPLICANT: Comoros, Timothy D.  
APPLICANT: Buttr, Timothy C.  
APPLICANT: Szymanski, Igor  
TITLE OF INVENTION: A LONG QT SYNDROME GENE  
FILE REFERENCE: 2123-113

CURRENT APPLICATION NUMBER: US/09/597,732

PRIOR FILING DATE: 2000-06-19

PRIOR APPLICATION NUMBER: 09/135,010

PRIOR FILING DATE: 1998-08-17

PRIOR APPLICATION NUMBER: 66/094,477

PRIOR FILING DATE: 1998-07-29

PRIOR APPLICATION NUMBER: 08/921,068

PRIOR FILING DATE: 1997-08-29

PRIOR APPLICATION NUMBER: 08/739,383

PRIOR FILING DATE: 1996-10-29

PRIOR APPLICATION NUMBER: 60/019,014

PRIOR FILING DATE: 1995-12-22

NUMBER OF SEQ ID NOS: 116

SEQUENCE: PatentIn Ver. 2.0

SEQ ID NO 4

LENGTH: 129

TYPE: PRT

ORGANISM: Homo sapiens

US-09-597-732-4

Query Match 21.0% Score 132.5; DB 4; Length 129;  
Best Local Similarity 45.1% Pred. No. 9.3e-08;  
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

OY 51 LYLAWMGESFVIALVSTVSKRRRHSNDPYQIYVED-WQERYKSQI 100  
DB 45 LYLAWLGFGEFTLGLIMLSTIRSKRLHSNDPNVYTESDAMQEKQNAV 95

RESULT 10  
US-09-679-185-2

Sequence 2, Application US/09679185

Patent No. 6458542

GENERAL INFORMATION:

INVENTOR: Robert, Daniel L.

TITLE OF INVENTION: METHOD OF SCREENING FOR SUSCEPTIBILITY TO

FILE REFERENCE: ATTORNEY DOCKET NO. 6458542 1242-33-2

CURRENT APPLICATION NUMBER: US/09/679,185

PRIOR FILING DATE: 1999-10-08

NUMBER OF SEQ ID NOS: 11

SEQUENCE: PatentIn Ver. 2.0

SEQ ID NO 2

LENGTH: 129

TYPE: PRT  
ORGANISM: Homo sapiens  
US-09-679-185-2

Query Match 21.0% Score 132.5; DB 4; Length 129;  
Best Local Similarity 45.1% Pred. No. 9.3e-08;  
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

OY 51 LYLAWMGESFVIALVSTVSKRRRHSNDPYQIYVED-WQERYKSQI 100  
DB 45 LYLAWLGFGEFTLGLIMLSTIRSKRLHSNDPNVYTESDAMQEKQNAV 95

RESULT 11  
US-08-118-101A-6

Sequence 6, Application US/08118101A

Patent No. 5620892

INVENTOR: Stephen E.

APPLICANT: Kruckeberg, Aron M.

TITLE OF INVENTION: A STRAIN OF SACCAROMYCES CEREVISIAE

TITLE OF INVENTION: EXPRESSING THE GENE ENCODING POTASSIUM TRANSPORTER MINK

NUMBER OF SEQUENCES: 16

CORRESPONDENCE ADDRESS:

STREET: P O Box 4000

CITY: Princeton

STATE: New Jersey

COUNTRY: U.S.A.

ZIP: 08543-4000

COMPUTER READABLE FORM:

MODIFIER TYPE: floppy disk

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/118,101A

FILING DATE: 435

CLASSIFICATION: A35

ATTORNEY/AGENT INFORMATION:

NAME: Gail, Timothy J.

REGISTRATION NUMBER: 33,111

TELEPHONE: (609) 252-5501

TELEFAX: (609) 252-4526

IMPORANT FOR SEQUENCE CHARACTERISTICS:

SEQUENCE: amino acid

LENGTH: 132 amino acids

TYPE: amino acid

MOLECULE TYPE: protein

US-08-118-101A-6

Query Match 21.0% Score 132.5; DB 1; Length 132;  
Best Local Similarity 45.1% Pred. No. 9.6e-08;  
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

OY 51 LYLAWMGESFVIALVSTVSKRRRHSNDPYQIYVED-WQERYKSQI 100  
DB 48 LYLAWLGFGEFTLGLIMLSTIRSKRLHSNDPNVYTESDAMQEKQNAV 98

RESULT 12  
US-09-069-896-4

Sequence 4, Application US/09069896

Patent No. 6071120

GENERAL INFORMATION:

INVENTOR: Hillman, Jennifer L.

APPLICANT: Celleron, Chandra

TITLE OF INVENTION: DELAYED RECTIFIER POTASSIUM

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01 COMPUTER: IBM Compatible
02 OPERATING SYSTEM: DOS
03 SOFTWARE: FASTSD for Windows Version 2.0
04 CURRENT APPLICATION DATA:
05 APPLICATION NUMBER: US/09/471,468
06 FILING DATE:
07 PRIORITY INFORMATION:
08 PRIOR APPLICATION DATA:
09 APPLICATION NUMBER: 09/069,896
10 FILING DATE:
11 ATTORNEY/AGENT INFORMATION:
12 NAME: Cerrione, Michael C
13 REGISTRATION NUMBER: 39,132
14 REFERENCE/DOCKET NUMBER: PP-0507 US
15 TELECOMMUNICATION INFORMATION:
16 TELEPHONE: 650-5510555
17 TELEFAX: 650-845-4160
18
19 INFORMATION FOR SEQ ID NO: 4:
20 SEQUENCE CHARACTERISTICS:
21 LENGTH: 130 amino acids
22 TYPE: amino acid
23 STRANDEDNESS: single
24 TOPOLOGY: linear
25 IMMEDIATE SOURCE:
26 ORGANISM: Oryzomys
27 CLONE: 203977
28
29 US-09-471-468-4
30
31 Query Match 20.3%; Score 128; DB 4; Length 130;
32 Best Local Similarity 41.4%; Pred. No. 3e-07;
33 Matches 29; Conservative 17; Mismatches 20; Indels 4; Gaps 3.
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35 Db 51 LTVLWMIQMSFVIVLVSVYSKRRHSNDYQIYED-0QERYS-0QIMESRSK 107
36 46 LTVLWMLDPPGFFGLDMLSTLSKSLHSDHIVESNDKRGKGLQATVLESFR 105
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38 Db 108 AT-FHENIG 116
39 106 ACTYENQAA 115
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41 RESULT 14
42 US-09-679-185-4
43 Sequence 4; Application US/09679185
44 Patent No. 6458542
45 GENERAL INFORMATION:
46 APPLICANT: George Jr., Alfred L.
47 APPLICANT: Roden, Dan M
48 TITLE OF INVENTION: METHOD OF SCREENING FOR SUSCEPTIBILITY TO
49 TITLE OF INVENTION: DRUG-INDUCED CARDIAC ARRYTHMIA
50 FILE REFERENCE: Attorney Docket No. 6458542 1242-33-2
51 CURRENT APPLICATION NUMBER: US/09/679,185
52 PRIORITY INFORMATION:
53 PRIOR APPLICATION NUMBER: 60/158,696
54 PRIOR FILING DATE: 1999-10-08
55 NUMBER OF SEQ ID NOS: 11
56 SOFTWARE: PatentIn Ver. 2.0
57
58 SEQ ID NO 4
59 LENGTH: 129
60 TYPE: PPT
61 ORGANISM: Homo sapiens
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63 US-09-679-185-4
64
65 Query Match 20.2%; Score 127.5; DB 4; Length 129;
66 Best Local Similarity 43.1%; Pred. No. 3.4e-07;
67 Matches 22; Conservative 16; Mismatches 12; Indels 1; Gaps 1.
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69 Db 51 LTVLWMIQMSFVIVLVSVYSKRRHSNDYQIYED-0QERYSQI 100
70 LTVLWMLDPPGFFGLDMLSTLSKSLHSDHIVESNDKRGKGLQATVLESFR 105
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72 Db 45 LTVLWMLDPPGFFGLDMLSTLSKSLHSDHIVESNDKRGKGLQATVLESFR 105

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RESULT 15
US-09-134-001C-5532
? Sequence 5532, Application US/09134001C
? Patent No. 6380370
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? GENERAL INFORMATION:
? APPLICANT: Lynn Doucette-Stamm et al
? TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO STAPHYLOCOCCUS
? TITLE OF INVENTION: EPIDERMIDIS FOR DIAGNOSTICS AND THERAPEUTICS
? FILING DATE: 1997-08-13
? CURRENT APPLICATION NUMBER: US/09/134,001C
? PRIOR FILING DATE: 1998-08-13
? PRIOR APPLICATION NUMBER: US 60/064, 964
? PRIOR FILING DATE: 1997-11-08
? PRIOR APPLICATION NUMBER: US 60/055, 779
? PRIOR FILING DATE: 1997-08-14
? NUMBER OF SEQ ID NOS: 5674
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? LENGTH: 262
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? TYPE: PRT
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? ORGANISM: Staphylococcus epidermidis
? US-09-134-001C-5532

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GenCore version 5.1.4.P5.4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 20:27:37 : Search time 100.392 Seconds

(Without alignments) 9628.006 Million cell updates/sec

File: US-09-550-163-1

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Searched: 828747 seqs, 660231138 residues

Total number of hits satisfying chosen parameters: 1657494

Minimum DB seq length: 0

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Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the total score distribution, and is derived by analysis of the score distribution.

## SUMMARIES

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4	372	50.8	372	10	US-09-864-761-31319		Sequence 31319, App
5	372	50.8	372	10	US-09-864-761-31319		Sequence 31319, App
6	372	50.8	372	10	US-09-864-761-31319		Sequence 31319, App
7	306	41.8	41.8	10	US-09-864-761-1671		Sequence 1671, App
8	306	41.8	41.8	10	US-09-864-761-1671		Sequence 1671, App
9	306	41.8	41.8	10	US-09-864-761-1671		Sequence 1671, App
10	53.2	7.3	390	10	US-09-184-316-3		Sequence 17593, App
11	53.2	7.3	1703	9	US-10-227-195A-1		Sequence 3, App1
12	53.2	7.3	113604	9	US-10-227-195A-2		Sequence 2, App1
13	50.8	6.3	368	10	US-09-864-761-4026		Sequence 4026, App
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15	36	4.9	1146	10	US-09-853-386-111		Sequence 111, App
16	36	4.9	1206	10	US-09-853-386-117		Sequence 117, App
17	36	4.9	1206	10	US-09-853-386-117		Sequence 117, App
18	36	4.9	1495	10	US-09-191-724-1		Sequence 1, App1
19	36	4.9	1495	10	US-09-191-724-1		Sequence 1, App1
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20	36	4.9	2579	10	US-09-823-830A-30		Sequence 30, App1
21	35.8	4.9	4057	9	US-10-239-676-181		Sequence 181, App
22	34.8	4.8	527	9	US-10-184-644-522		Sequence 522, App
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26	33.6	4.6	653	9	US-10-184-644-402		Sequence 402, App
27	33.6	4.6	1140	12	US-10-003-356-6		Sequence 6, App1
28	33.6	4.6	2781	12	US-10-003-356-9		Sequence 9, App1
29	33.4	4.6	4104	9	US-09-992-598-277		Sequence 277, App
30	33.4	4.6	4104	9	US-09-989-253A-277		Sequence 277, App
31	33.4	4.6	4104	9	US-09-989-253A-277		Sequence 277, App
32	33.4	4.6	4104	9	US-09-989-253A-277		Sequence 277, App
33	33.4	4.6	4104	9	US-09-989-253A-277		Sequence 277, App
34	33.4	4.6	4104	9	US-09-989-253A-277		Sequence 277, App
35	33.4	4.6	4104	9	US-09-989-253A-277		Sequence 277, App
36	33.4	4.6	4104	9	US-09-989-253A-277		Sequence 277, App
37	33.4	4.6	4104	9	US-09-989-253A-277		Sequence 277, App
38	33.4	4.6	4104	9	US-09-989-253A-277		Sequence 277, App
39	33.4	4.6	4104	9	US-09-989-253A-277		Sequence 277, App
40	33.4	4.6	4104	9	US-09-989-253A-277		Sequence 277, App
41	33.4	4.6	4104	9	US-09-989-253A-277		Sequence 277, App
42	33.4	4.6	4104	9	US-10-121-049-449		Sequence 449, App
43	33.4	4.6	4104	9	US-10-121-049-449		Sequence 449, App
44	33.4	4.6	4104	9	US-10-121-049-449		Sequence 449, App
45	33.4	4.6	4104	9	US-10-121-049-449		Sequence 449, App

## ALIGNMENTS

RESULT 1  
US-10-000-151B-5  
Sequence 5, Application US/10000151B  
Publication No. US2003013136A1  
GENERAL INFORMATION: US/10000151B  
APPLICANT: Basset, Jeffrey R.  
TITLE OF INVENTION: HUMAN KCR1 REGULATION OF HERG POTASSIUM CHANNEL BLOCK  
CURRENT APPLICATION NUMBER: US/10/000.151B  
CURRENT FILING DATE: 2000-10-30  
NUMBER OF SEQ ID NOS: 5  
SOFTWARE: Patent version 3.1  
SEQ ID NOS: 1-5  
LENGTH: 732  
ORGANISM: Homo sapiens  
US-10-000-151B-5

Query Match 99.8% Score 730.4 : DB 9 : Length 732:  
Best Local Similarity 99.9% : Pred. No. 0.0e+215 :  
Matches 731 : Mismatches 1 : Indels 0 : Gaps 0 :

QY	1	CAATTCGAAAGATCGCTTTTACGCTTCCCATTTTATTAATTCGA	60
DB	1	CAATTCGAAAGATCGCTTTTACGCTTCCCATTTTATTAATTCGA	60
QY	61	CGAGAGGAGACATCTGCTTATTCATATTCACAGACGCTTGACGCTTCG	120
DB	61	CGAGAGGAGACATCTGCTTATTCATATTCACAGACGCTTGACGCTTCG	120
QY	121	AAGATTTTATTCATATTCATATTCATATTCATATTCATATTCATATTCAT	180
DB	121	AAGATTTTATTCATATTCATATTCATATTCATATTCATATTCATATTCAT	180
QY	181	CTTCAGACCAAGTATCTGAGACATCTGATATTCATATTCATATTCATATTCAT	240
DB	181	CTTCAGACCAAGTATCTGAGACATCTGATATTCATATTCATATTCATATTCAT	240
QY	241	GATGGAATGCTTCATATTCATATTCATATTCATATTCATATTCATATTCAT	300
DB	241	GATGGAATGCTTCATATTCATATTCATATTCATATTCATATTCATATTCAT	300

```

Oy 301 ACGGAAACACCTCCATGACCCCTACACACAGATGATTTGAGAGACGTGGCGGAAAGATG 360
Db 301 ACGGAAACACCTCCATGACCCCTACACACAGATGATTTGAGAGACGTGGCGGAAAGATG 360
Oy 361 CAAAGGCGCAATCTGAAATCTGAGAGATGAGAGGCGACACATCTGATAGAAACATGGTGC 420
Db 361 CAAAGGCGCAATCTGAAATCTGAGAGATGAGAGGCGACACATCTGATAGAAACATGGTGC 420
Oy 421 GCGTGGGTTCAAAATGTGCTCCCTGATAGAGGAGAGGACACACAGCTAACATCTGACATC 480
Db 421 GCGTGGGTTCAAAATGTGCTCCCTGATAGAGGAGAGGACACACAGCTAACATCTGACATC 480
Oy 481 CAGACATGAGAGATGCGCAGTGCACAGGAGGCAATTCCTTCTGCTTGGAGAA 540
Db 481 CAGACATGAGAGATGCGCAGTGCACAGGAGGCAATTCCTTCTGCTTGGAGAA 540
Oy 541 GTGAGTCTCTGCTCTGCTGATGAGATTTTCATGAGATTTGAGTGGTGGCATMAAGA 600
Db 541 GTGAGTCTCTGCTCTGCTGATGAGATTTTCATGAGATTTGAGTGGTGGCATMAAGA 600
Oy 601 TGAATGACATTTCAATCTCAGATGATTTATGCTGCTGTGGAGCAATATTGGTGTGA 660
Db 601 TGAATGACATTTCAATCTCAGATGATTTATGCTGCTGTGGAGCAATATTGGTGTGA 660
Oy 661 AAGCCTCTTTACTTCCGGGCAAGATGATGATTTATCAATATCAATGATGCAAAAT 720
Db 661 AAGCCTCTTTACTTCCGGGCAAGATGATGATTTATCAATATCAATGATGCAAAAT 720
Oy 721 AAGCCCAATTT 732
Db 721 AAGCCCAATTT 732

```

```

RESULT 2
US-10-227-195A-1
: Sequence 1, Application US/10227195A
: Publication No. US2003007633A1
: GENERAL INFORMATION:
: APPLICANT: David
: APPLICANT: Arnold Deana
: TITLE OF INVENTION: Haplotype structure of chromosome 21
: FILE REFERENCE: 103001
: CURRENT APPLICATION NUMBER: US/10/227.195A
: CURRENT FILING DATE: 2002-11-18
: NUMBER OF SEQ ID NOS: 2
: ORIGIN: FastaSeq for Windows Version 4.0
: SEQ ID NO: 1
: LENGTH: 113604
: TYPE: DNA
: ORGANISM: Human
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: 7175..7204, 36973..66372, 76921..81512, 88727
: OTHER FEATURES:
: SOFTWARE: FastaSeq for Windows Version 4.0
US-10-227-195A-1

```

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Query Match 99.88; Score 730.4; DB 9; Length 113604;
Best Local Similarity 99.98; Pred. No. 1.5e-213;
Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

```

```

Oy 1 CAAATCCGAAAGAACGCTTTCTCAACCTTGCGCTATTTATATTAATTAATGCA 60
Db 17403 CAAATCCGAAAGAACGCTTTCTCAACCTTGCGCTATTTATATTAATTAATGCA 17462
Oy 61 GCAGAGGAGGAAGCATGTCTACTTTATCAATTTCAACAGACGTGGAAAGCGTCTCG 120
Db 17463 GCAGAGGAGGAAGCATGTCTACTTTATCAATTTCAACAGACGTGGAAAGCGTCTCG 17522
Oy 121 AAGGATTTTATCTTATATGACAAATTTGGCGGCAGAACACAAAGCTGAGCAAGGC 180
Db 17523 AAGGATTTTATCTTATATGACAAATTTGGCGGCAGAACACAAAGCTGAGCAAGGC 17582

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Oy 181 CCTCCAGACCAAGGTGATGCTGAGAACTTCACTATGATCTGACATGATGAT 240
Db 17583 CCTCCAGACCAAGGTGATGCTGAGAACTTCACTATGATCTGACATGATGAT 17642
Oy 241 GATTGGAAATGCTCTTCTGATCAATGCTGGACATCTGGGAGACATCCGAAT 300
Db 17643 GATTGGAAATGCTCTTCTGATCAATGCTGGACATCTGGGAGACATCCGAAT 17702
Oy 301 ACGGGAACATCCATGACCCCTACACACAGTANATGTAGAGATCGGCGGAAAAAGTA 360
Db 17703 ACGGGAACATCCATGACCCCTACACACAGTANATGTAGAGATCGGCGGAAAAAGTA 17762
Oy 361 CAAAGCGCAATCTGAAATCTGAGAGATGAGAGGCGACACATCTGAGAAACATGGTGC 420
Db 17763 CAAAGCGCAATCTGAAATCTGAGAGATGAGAGGCGACACATCTGAGAAACATGGTGC 17822
Oy 421 GCGTGGGTTCAAAATGTGCTCCCTGATAGAGGAGAGGACACACAGCTAACATCTGACATC 480
Db 17823 GCGTGGGTTCAAAATGTGCTCCCTGATAGAGGAGAGGACACACAGCTAACATCTGACATC 17882
Oy 481 CAGACATGAGAGATGCGCAGTGCACAGGCAATTCCTTCTGCTTGGAGAA 540
Db 17883 CAGACATGAGAGATGCGCAGTGCACAGGCAATTCCTTCTGCTTGGAGAA 17942
Oy 541 GTGAGTCTCTGCTCTGCTGATGAGATTTTCATGAGATTTGAGTGGTGGCATMAAGA 600
Db 17943 GTGAGTCTCTGCTCTGCTGATGAGATTTTCATGAGATTTGAGTGGTGGCATMAAGA 18002
Oy 601 TGAATGACATTTCAATCTCAGATGATTTATGCTGCTGTGGAGCAATATTGGTGTGA 660
Db 18003 TGAATGACATTTCAATCTCAGATGATTTATGCTGCTGTGGAGCAATATTGGTGTGA 18062
Oy 661 AAGCCTCTTTACTTCCGGGCAAGATGATGATTTATCAATATCAATGATGCAAAAT 720
Db 18063 AAGCCTCTTTACTTCCGGGCAAGATGATGATTTATCAATATCAATGATGCAAAAT 18122
Oy 721 AAGCCCAATTT 732
Db 18123 AAGCCCAATTT 18134

```

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RESULT 3
US-10-227-195A-2
: Sequence 2, Application US/10227195A
: Publication No. US2003007633A1
: GENERAL INFORMATION:
: APPLICANT: Cox, David
: APPLICANT: Arnold, Deana
: TITLE OF INVENTION: Haplotype structure of chromosome 21
: FILE REFERENCE: 103001
: CURRENT APPLICATION NUMBER: US/10/227.195A
: CURRENT FILING DATE: 2002-11-18
: NUMBER OF SEQ ID NOS: 2
: ORIGIN: FastaSeq for Windows Version 4.0
: SEQ ID NO: 2
: LENGTH: 113604
: TYPE: DNA
: ORGANISM: Human
US-10-227-195A-2

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Query Match 99.88; Score 730.4; DB 9; Length 113604;
Best Local Similarity 99.98; Pred. No. 1.5e-213;
Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

```

```

Oy 1 CAAATCCGAAAGAACGCTTTCTCAACCTTGCGCTATTTATATTAATTAATGCA 60
Db 17403 CAAATCCGAAAGAACGCTTTCTCAACCTTGCGCTATTTATATTAATTAATGCA 17462
Oy 61 GCAGAGGAGGAAGCATGTCTACTTTATCAATTTCAACAGACGTGGAAAGCGTCTCG 120
Db 17463 GCAGAGGAGGAAGCATGTCTACTTTATCAATTTCAACAGACGTGGAAAGCGTCTCG 17522
Oy 121 AAGGATTTTATCTTATATGACAAATTTGGCGGCAGAACACAAAGCTGAGCAAGGC 180

```



```

GENERAL INFORMATION:
APPLICANT: Penn, Sharon G.
APPLICANT: Hanzel, David K.
APPLICANT: Chen, Wensheng
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
FILE REFERENCE: Aeomlca-x-1
CURRENT APPLICATION NUMBER: US/09/864,761
PRIOR APPLICATION NUMBER: US 60/180,312
PRIOR FILING DATE: 2000-02-04
PRIOR APPLICATION NUMBER: US 60/207,456
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: US 09/632,366
PRIOR FILING DATE: 2000-08-03
PRIOR APPLICATION NUMBER: US 60/236,359
PRIOR FILING DATE: 2000-10-04
PRIOR APPLICATION NUMBER: US 60/236,359
PRIOR FILING DATE: 2000-09-27
PRIOR APPLICATION NUMBER: PCT/US01/00666
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00667
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00664
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00669
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00665
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00668
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00663
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00662
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00661
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00670
PRIOR FILING DATE: 2000-09-21
PRIOR APPLICATION NUMBER: US 60/234,687
PRIOR FILING DATE: 2000-06-30
PRIOR APPLICATION NUMBER: US 09/608,408
PRIOR FILING DATE: 2001-01-29
PRIOR APPLICATION NUMBER: US 09/774,203
SOFTWARE: ABLINK Sequence Listing Engine vers. 1.1
SEQ ID NO 3463
LENGTH: 450
TYPE: DNA
ORGANISM: Homo sapiens
FEATURES:
OTHER INFORMATION: MAP TO AP000052.1 MARROW SIGNAL - 1.1
OTHER INFORMATION: EXPRESSED IN BILLO SIGNAL - 1.1
OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL - 1.3
OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL - 0.92
OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL - 1.4
OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL - 1
OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL - 1.1
OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL - 1.2
OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL - 0.94
OTHER INFORMATION: EXPRESSED IN BILLY, SIGNAL - 0.88
US-09-864-761.3463
Query Match 50.3%; Score 368; DB 10; Length 450;
Best Local Similarity 100.0%; Pred. No. 3.3e-103;
Matches 368; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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61 GCAGAGGAGAGACATCTACTTATTCATTTTACACAGCCGTGAGACGCTTCG 120
143 GACAGAGACAGACATCTACTTATTCATTTTACACAGCCGTGAGACGCTTCG 202
121 AAGGATTTTATTTACTTATTAATGACATATTCGCGCGAGACACAGCTGACGAGAGG 180
203 AAGGATTTTATTTACTTATTAATGACATATTCGCGCGAGACACAGCTGACGAGAGG 262
181 CCTCAGACCCAGATGTATGTATGATACATCTACTTATTCATTTTACACAGCTTCGAT 240
263 CTTCCAGACCCAGATGTATGTATGATACATCTACTTATTCATTTTACACAGCTTCGAT 322
241 GATTGAGATGTCTCTTATCATCATGCGCATCTCGTGACACATGTAATCCAGAG 300
323 ATTTGAGATGTCTCTTATCATCATGCGCATCTCGTGACACATGTAATCCAGAG 382
301 AGCGGAGACATGAGAGACCGCTGACACATCTACTTATTCATTTTACACAGCTTCGAT 360
383 AGCGGAGACATGAGAGACCGCTGACACATCTACTTATTCATTTTACACAGCTTCGAT 442
361 CAGAGACC 368
443 CAGAGACC 450
RESULT 6
US-09-864-761-20233
Sequence 20233, Application US/09864761
Patent No. US20020048763A1
GENERAL INFORMATION:
APPLICANT: Penn, Sharon G.
APPLICANT: Hanzel, David K.
APPLICANT: Chen, Wensheng
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
FILE REFERENCE: Aeomlca-x-1
CURRENT APPLICATION NUMBER: US/09/864,761
PRIOR FILING DATE: 2001-05-23
PRIOR FILING DATE: 2000-02-04
PRIOR FILING DATE: 2000-02-04
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: US 09/632,366
PRIOR FILING DATE: 2000-08-03
PRIOR APPLICATION NUMBER: US 09/632,366
PRIOR FILING DATE: 2000-09-27
PRIOR APPLICATION NUMBER: PCT/US01/00666
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00667
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00664
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00669
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00665
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00668
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00663
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00662
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00661
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: US 60/234,687
PRIOR FILING DATE: 2000-09-21
PRIOR APPLICATION NUMBER: US 09/608,408
PRIOR APPLICATION NUMBER: US 09/774,203

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	PRIOR APPLICATION NUMBER:	US 09/632,366
	PRIOR FILING DATE:	2000-08-03
	PRIOR APPLICATION NUMBER:	GB 24263, 6
	PRIOR FILING DATE:	2000-10-04
	PRIOR APPLICATION NUMBER:	US 60/236,359
	PRIOR FILING DATE:	2000-09-27
	PRIOR APPLICATION NUMBER:	ECT/US01/00666
	PRIOR FILING DATE:	2001-01-30
	PRIOR APPLICATION NUMBER:	ECT/US01/00667
	PRIOR FILING DATE:	2001-01-30
	PRIOR APPLICATION NUMBER:	ECT/US01/00664
	PRIOR FILING DATE:	2001-01-30
	PRIOR APPLICATION NUMBER:	ECT/US01/00669
	PRIOR FILING DATE:	2001-01-30
	PRIOR APPLICATION NUMBER:	ECT/US01/00665
	PRIOR FILING DATE:	2001-01-30
	PRIOR APPLICATION NUMBER:	ECT/US01/00668
	PRIOR FILING DATE:	2001-01-30
	PRIOR APPLICATION NUMBER:	ECT/US01/00663
	PRIOR FILING DATE:	2001-01-30
	PRIOR APPLICATION NUMBER:	ECT/US01/00662
	PRIOR FILING DATE:	2001-01-30
	PRIOR APPLICATION NUMBER:	ECT/US01/00661
	PRIOR FILING DATE:	2001-01-30
	PRIOR APPLICATION NUMBER:	ECT/US01/00670
	PRIOR FILING DATE:	2001-01-30
	PRIOR FILING DATE:	2000-09-22
	PRIOR FILING DATE:	2000-06-30
	PRIOR FILING DATE:	2000-06-30
	PRIOR APPLICATION NUMBER:	US 09/774,203
	PRIOR FILING DATE:	2001-01-29
	NUMBER OF SEQ ID NOS:	49117
	SOFTWARE:	Anomax Sequence Listing Engine vstrs. 1.1
	SEQ ID NO 16671	
	LENGTH:	471
	TYPE:	DNA
	ORGANISM:	Homo sapiens
	FEATURE:	
	OTHER INFORMATION:	MAP TO AP000120.1
	OTHER INFORMATION:	EXPRESSED IN HELA, SIGNAL = 0.98
	OTHER INFORMATION:	EXPRESSED IN HBL100, SIGNAL = 0.67
	US-09-864-761-16671	
	Query Match	41 8%: Score 306; DB 10; Length 471;
	Best Local Similarity	100.0%; Pval No N 4.6e-84;
	Matches 306; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
Gy	1	CAATTCAGAAANAATCCGTTTCTTAACGTGGTCCCATATTATTTAATAATGCA 60
Gy	166	CAAATCGAGAAMATCAGCTTTTCCAACTTGCCATATTTATTAATAATGCA 225
Oy	61	GCGAGGAGAACACTGCCTATTTCCAACTTGCCATATTTATTAATAATGCA 120
Dy	226	CGAGGAGGAACACTGCCTACTTTTCATATTTCCACAAGAGCTGGAACGCTTCCG 285
Oy	121	AAGATTTTTATTTACTTATATGAGCAATATGGGCCGACAGCAACAAGTAGTACAAAGGC 180
Dy	286	AAGATTTTTATTTACTTATATGAGCAATATGGGCCGACAGCAACAAGTAGTACAAAGGC 345
Oy	181	CGTCACGCAAAATGTATATGAGCAATATGGGCCGACAGCAACAAGTAGTACAAAGGC 240
Dy	346	CGTCACGCAAAATGTATATGAGCAATATGGGCCGACAGCAACAAGTAGTACAAAGGC 405
Oy	241	GATTGGAAATTCCTCTTCATCATCGTGGCCATCCGTGGTGAACAACGTGAATCAAGG 300
Dy	406	GATTGGAAATTCCTCTTCATCATCGTGGCCATCCGTGGTGAACAACGTGAATCAAGG 465
Oy	301	
Dy	466	ACGGGA 471



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? OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 3.6
? OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 3.9
? OTHER INFORMATION: EXPRESSED IN BT47A, SIGNAL = 2.4
? OTHER INFORMATION: EXPRESSED IN BOVE MARROW, SIGNAL = 1.8
? OTHER INFORMATION: EXPRESSED IN HB1100, SIGNAL = 1.8
? OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 2.8
? OTHER INFORMATION: EXPRESSED IN SPLEEN, SIGNAL = 1.8
? OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 2.3
? OTHER INFORMATION: EXPRESSED IN LONG, SIGNAL = 3.2
? OTHER INFORMATION: EST_HUMAN HIT: AA70188.1, EVALU6 6.30e-02
? OTHER INFORMATION: SWISSPROT HIT: P15382, EVALU6 4.00e-61
? OTHER INFORMATION: NT HIT: g11156222, EVALU6 0.00e+00
US-08-864-761-17593

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Query Match 7.3%; Score 53.2; DB 10; Length 390;

Best Local Similarity 63.6%; Pred. No. 4.2e-05; Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;

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Oy 223 CCTGTAACCTCATGTGATGATGGAATGTTCTCTTCATCATCGTCGCACTCGTGAG 282
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 209 CCTGTAACCTCATGTGATGATGGAATGTTCTCTTCATCATCGTCGCACTCGTGAG 150
Oy 283 CACGTGAAATTCAGAGAGCGGAACCTCCATGATGACCCCTACACAGCAATGTTG--T 339
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 149 CTACATCCGCTCCAGAGCTGGAGACCTCCAGACACCCTTCAACGCTCATCATGAGTGC 90
Oy 340 AGAGACTGCGCAGAAAGATGACAGAGCCCAATC 373
Db 89 CATGCTCGTCAGAGAGAGAGACAGAGCCCTATGTC 56

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RESULT 10

US-10-138-316-3  
Sequence 3, Application US/10138316  
Publication No. US20030054380A1

GENERAL INFORMATION: Mark T.  
APPLICANT: Sanofi-Schering  
APPLICANT: Sanofi-Schering, Michael C.

TITLE OF INVENTION: MUTATIONS IN THE KCNEL GENE ENCODING HUMAN MARK WHICH  
TITLE OF INVENTION: CAUSE ARRYTHMIA SUSCEPTIBILITY THEREBY ESTABLISHING

FILE REFERENCE: 733-162 US/0-218, 316

CURRENT FILING DATE: 2002-05-06

PRIOR APPLICATION NUMBER: 09/444,295

PRIOR FILING DATE: 1999-11-22

PRIOR APPLICATION NUMBER: 09/135,020

PRIOR FILING DATE: 1998-08-17

PRIOR APPLICATION NUMBER: 08/921,068

PRIOR FILING DATE: 1997-08-29

PRIOR APPLICATION NUMBER: 08/739,383

PRIOR FILING DATE: 1996-10-29

PRIOR APPLICATION NUMBER: 60/019,014

PRIOR FILING DATE: 1995-12-22

PRIOR APPLICATION NUMBER: 60/094,477

PRIOR FILING DATE: 1998-07-29

NUMBER OF SEQ ID NOS: 114

SOFTWARE: patent ver. 2.0

SEQ ID NO 3

LENGTH: 1703

TYPE: DNA

ORGANISM: Homo sapiens

NAME/KEY: CDS

LOCATION: (193)..(579)

NAME/KEY: misc

LOCATION: (1)..(1703)

OTHER INFORMATION: n may be any nucleotide

US-10-138-316-3

Query Match 7.3%; Score 53.2; DB 9; Length 1703;

Best Local Similarity 63.6%; Pred. No. 1e-05; Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;

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Oy 223 CCTGTAACCTCATGTGATGATGGAATGTTCTCTTCATCATCGTCGCACTCGTGAG 282
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 324 CCTGTAACCTCATGTGATGATGGAATGTTCTCTTCATCATCGTCGCACTCGTGAG 383
Oy 283 CACGTGAAATTCAGAGAGCGGAACCTCCATGATGACCCCTACACAGCAATGTTG--T 339
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 384 CTACATCCGCTCCAGAGCTGGAGACCTCCAGACACCCTTCAACGCTCATCATGAGTGC 443
Oy 340 AGAGACTGCGCAGAAAGATGACAGAGCCCAATC 373
Db 444 CATGCTCGTCAGAGAGAGAGACAGAGCCCTATGTC 477

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RESULT 11

US-10-227-195A-2/C  
Sequence 1, Application US/10227195A  
Publication No. US2003007633A1

GENERAL INFORMATION: Arnold, Deana  
APPLICANT: Arnold, Deana

TITLE OF INVENTION: Haplotype structure of chromosome 21

FILE REFERENCE: 103001

CURRENT FILING DATE: 2002-11-18

CURRENT APPLICATION NUMBER: US/10/227,195A

NUMBER OF SEQ ID NOS: 2

SOFTWARE: FASTEST for Windows Version 4.0

SEQ ID NO

LENGTH: 113604

TYPE: DNA

ORGANISM: Human

NAME/KEY: misc:feature

LOCATION: 7175, 7204, 36973, 76921, 81512, 88727

OTHER INFORMATION: n - G or C

US-10-227-195A-1

Query Match 7.3%; Score 53.2; DB 9; Length 113604;

Best Local Similarity 63.6%; Pred. No. 0.00014;

Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;

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Oy 223 CCTGTAACCTCATGTGATGATGGAATGTTCTCTTCATCATCGTCGCACTCGTGAG 282
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 96499 CCTGTAACCTCATGTGATGATGGAATGTTCTCTTCATCATCGTCGCACTCGTGAG 96440
Oy 283 CACGTGAAATTCAGAGAGCGGAACCTCCATGATGACCCCTACACAGCAATGTTG--T 339
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 96439 CTACATCCGCTCCAGAGCTGGAGACCTCCAGACACCCTTCAACGCTCATCATGAGTGC 96380
Oy 340 AGAGACTGCGCAGAAAGATGACAGAGCCCAATC 373
Db 96379 CATGCTCGTCAGAGAGAGAGACAGAGCCCTATGTC 96346

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RESULT 12

US-10-227-195A-2/C  
Sequence 2, Application US/10227195A  
Publication No. US2003007633A1

GENERAL INFORMATION: Arnold, Deana  
APPLICANT: Cox, David

TITLE OF INVENTION: Haplotype structure of chromosome 21

FILE REFERENCE: 103001

CURRENT FILING DATE: 2002-11-18

CURRENT APPLICATION NUMBER: US/10/227,195A

NUMBER OF SEQ ID NOS: 2

SOFTWARE: FASTEST for Windows Version 4.0

SEQ ID NO 2

LENGTH: 113604

TYPE: DNA

ORGANISM: Human



:/ PRIOR FILING DATE: 2001-01-30  
:/ PRIOR APPLICATION NUMBER: US 60/234,687  
:/ PRIOR FILING DATE: 2000-09-21  
:/ PRIOR APPLICATION NUMBER: US 09/608,408  
:/ PRIOR FILING DATE: 2000-06-30  
:/ PRIOR APPLICATION NUMBER: US 09/774,203  
:/ PRIOR FILING DATE: 2001-01-29  
:/ NUMBER OF SEQ ID NOS: 49117  
:/ SOFTWARE: Anomax Sequence Listing Engine vers. 1.1  
:/ SEQ ID NO 810  
:/ LENGTH: 381  
:/ TYPE: DNA  
:/ ORGANISM: Homo sapiens  
:/ PROVENANCE: 1  
:/ OTHER INFORMATION: MAP TO AP000168.1  
:/ OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 2.6  
:/ OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 3.6  
:/ OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 3.9  
:/ OTHER INFORMATION: EXPRESSED IN BT474, SIGNAL = 2.4  
:/ OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.8  
:/ OTHER INFORMATION: EXPRESSED IN HEL100, SIGNAL = 1.8  
:/ OTHER INFORMATION: EXPRESSED IN HEL100, SIGNAL = 2.0  
:/ OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1.8  
:/ OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 2.3  
:/ OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 3.2  
US-09-864-761-810

## Query Match

Best Local Similarity 6.38; Score 46; DB 10; Length 381;  
Best Local Similarity 64.28; Pred. No. 0.00069;  
Matches 86; Conservative 0; Mismatches 45; Indels 3; Gaps 1;

OY 243 TTGGAAATGTTCTTTCATCATCGGGCCATCGTGTGAGCAGCTGTGAATCCAGAGAGC 302  
DB 378 TGGGATTCCTGCTGCTTCACCTCGGCGCATCGTGTGAGCTACATCGCGCTCCAGAGAGC 319  
OY 303 GGGAGACCTCATGATGACCCCTACACAGCTACATGTGA--GAGAGTGGCAGGAAAGT 359  
DB 318 TGGAGCTCATGAGAGCCGCTCAAGCTGTACATGAGTCCATGATCGCTGCAAGAGAGG 259  
OY 360 ACAAGAGCCCAATC 373  
DB 258 ACAAGGCTTATGTC 245

## RESULT 15

US-09-853-386-111  
:/ Sequence 111, Application US/09853386  
:/ Patent No. US20020049151A1  
:/ GENERAL INFORMATION:  
:/ APPLICANT: Murphy, Evelyn  
:/ APPLICANT: Freshman, Barry  
:/ APPLICANT: Murphy, David  
:/ APPLICANT: Fitzgerald, Oliver  
:/ TITLE OF INVENTION: Therapeutic Approaches to Diseases by Suppression of the NTR  
:/ FILE REFERENCE: P01972051  
:/ CURRENT APPLICATION NUMBER: US/09/853,386  
:/ PRIOR FILING DATE: 2001-05-11  
:/ PRIOR APPLICATION NUMBER: US 60/203645  
:/ PRIOR FILING DATE: 2000-05-12  
:/ NUMBER OF SEQ ID NOS: 153  
:/ SOFTWARE: PatentIn version 3.1  
:/ SEQ ID NO 111  
:/ LENGTH: 1146  
:/ TYPE: DNA  
:/ ORGANISM: HUMAN  
:/ US-09-853-386-111

Query Match 4.98; Score 36; DB 10; Length 1146;  
Best Local Similarity 52.78; Pred. No. 1.7; Indels 0; Gaps 0;  
Matches 78; Conservative 0; Mismatches 0;

OY 188 GCCAAAGTGATGCTGAGACCTTCACTATGTCATCCCTGACTCATGCTGATGATTTGGA 247

DB 679 GCGAAAAGCCCTGGGGTGTACACCGACATCACTACAGGCCCCCATGATCTGTGCTGTG 738  
OY 248 ATGTTCTTTTCATCATGCTGTGACCATCTGCTGTGAGCAGCTGTGAATCCAGAGAGG 307  
DB 739 CTGATCAATTTCAATCTTCCTTTTCACATGCTGCTGCGCATCTGCTCATGACCAAGCTCG 798  
OY 308 CACTCAATGACCCCTCACACCAAGATACA 335  
DB 739 TCCACCACTGCTGAGACCACTCATGATACA 826

Search completed: May 21, 2003, 22:42:03  
Job time : 217.392 secs



GenCore version 5.1.4-P5.4578  
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OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 20:14:00 ; Search time 42.5504 Seconds  
(without alignments)

5275.799 Million cell updates/sec

Title: US-09-550-163-1  
Perfect score: 732  
Sequence: 1 caatccagaagaatcgcgt.....atgaataaagccaattt 732  
Scoring table: IDENTITY/MNC  
Gapop 10.0 ; Gapext 1.0

Searched: 441362 seqs, 1533381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :

- 1: /cgn2.6/p/odata/1/lna/5A.COMB.seq.\*
- 2: /cgn2.6/p/odata/1/lna/5B.COMB.seq.\*
- 3: /cgn2.6/p/odata/1/lna/6A.COMB.seq.\*
- 4: /cgn2.6/p/odata/1/lna/6B.COMB.seq.\*
- 5: /cgn2.6/p/odata/1/lna/PCUTS.COMB.seq.\*
- 6: /cgn2.6/p/odata/1/lna/Backfiles.seq.\*

Prod. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	53.2	7.3	398	4	US-08-118-101A-5	Sequence 5, Appl1
2	53.2	7.3	436	4	US-09-679-185-1	Sequence 1, Appl1
3	53.2	7.3	1703	3	US-09-135-021-77	Sequence 77, Appl1
4	53.2	7.3	1703	4	US-09-135-020-3	Sequence 3, Appl1
5	53.2	7.3	1703	4	US-09-135-010A-3	Sequence 3, Appl1
6	53.2	7.3	1703	4	US-09-444-871-3	Sequence 3, Appl1
7	53.2	7.3	1703	4	US-09-597-733-3	Sequence 3, Appl1
8	53.2	7.3	1703	4	US-09-597-732-3	Sequence 3, Appl1
9	53.2	7.3	1703	4	US-09-597-732-3	Sequence 3, Appl1
10	51.6	7.0	436	4	US-09-679-185-3	Sequence 3, Appl1
11	36.6	5.0	2653	1	US-08-318-831-1	Sequence 1, Appl1
12	36.2	4.9	7218	1	US-08-232-463-14	Sequence 14, Appl1
13	36	4.9	1380	1	US-08-110-286A-1	Sequence 1, Appl1
14	36	4.9	1380	1	US-08-482-745-14	Sequence 1, Appl1
15	36	4.9	1492	4	US-08-069-896-2	Sequence 2, Appl1
16	33.6	4.6	645	3	US-09-471-468-2	Sequence 2, Appl1
17	33	4.5	606	4	US-09-328-111-133	Sequence 133, Appl1
18	33	4.5	606	4	US-09-307-143-3	Sequence 3, Appl1
19	33	4.5	2912	4	US-08-986-416-303	Sequence 303, Appl1
20	32.4	4.4	837	4	US-09-221-017B-923	Sequence 923, Appl1
21	32.4	4.4	4659	4	US-08-724-394A-20	Sequence 1, Appl1
22	32.4	4.4	21630	3	US-08-724-394A-21	Sequence 20, Appl1
23	32	4.4	24640	2	US-08-724-394A-21	Sequence 21, Appl1
24	32	4.4	24640	2	US-08-724-394A-22	Sequence 22, Appl1
25	32	4.4	24640	2	US-08-724-394A-21	Sequence 22, Appl1
26	31.4	4.3	566	4	US-09-221-017B-919	Sequence 919, Appl1
27	30.6	4.2	744	3	US-08-965-644-17	Sequence 17, Appl1

28	30.6	4.2	744	3	US-08-444-189-17	Sequence 17, Appl
29	30.6	4.2	744	3	US-08-468-544-17	Sequence 17, Appl
30	30.6	4.2	2929	4	US-09-705-299-10	Sequence 10, Appl
31	30.6	4.2	6822	4	US-09-426-998-3	Sequence 3, Appl1
32	30.6	4.2	7502	3	US-08-969-644-6	Sequence 6, Appl1
33	30.6	4.2	7502	3	US-08-444-189-6	Sequence 6, Appl1
34	30.6	4.2	7741	4	US-09-426-998-4	Sequence 4, Appl1
35	30.6	4.2	98844	4	US-09-751-211-10	Sequence 10, Appl
36	30.6	4.2	28473	4	US-08-961-527-83	Sequence 83, Appl
37	30.2	4.1	28001	4	US-09-819-993-3	Sequence 3, Appl1
38	30.2	4.1	33000	4	US-09-215-694-18	Sequence 18, App
39	30.2	4.1	1001	4	US-09-641-638-198	Sequence 198, App
40	29.8	4.1	12266	4	US-08-724-394A-18	Sequence 18, Appl
41	29.8	4.1	72604	4	US-09-426-998-4	Sequence 4, Appl1
42	29.8	4.0	72604	4	US-09-426-998-4	Sequence 4, Appl1
43	29.6	4.0	462	3	US-08-863-813A-33	Sequence 33, Appl
44	29.6	4.0	3273	6	US-08-965-644-17	Sequence 17, Appl
45	29.6	4.0	3273	6	US-08-965-644-17	Sequence 17, Appl

## ALIGNMENTS

RESULT 1

US-08-118-101A-5  
Sequence 5, Application US/08118101A  
Patent No. 5620892

GENERAL INFORMATION:

APPLICANT: KUTZ, Stephen E.  
INVENTOR: KUTZ, Stephen E.  
TITLE OF INVENTION: A STRAIN OF SACHAROWYCES CREVITIAE

NUMBER OF SEQUENCES: 16  
CORRESPONDENCE ADDRESS:

ADDRESSER: Burton Rodney  
STREET: P.O. Box 4000  
CITY: New Jersey

COUNTRY: U.S.A.  
ZIP: 08543-4000

COMPUTER TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
SOFTWARE SYSTEM: PC DOS/MS-DOS

APPLICATION NUMBER: US/08/118.101A  
CURRENT APPLICATION DATA: Version #1.25

CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:

NAME: Gaul, Timothy J.  
REFERENCE/DOCKET NUMBER: DC227

TELECOMMUNICATION INFORMATION:  
TELEPHONE: (609) 252-5901  
TELEFAX: (609) 252-4526

INFORMATION FOR SEQ ID NO: 5:  
SEQUENCE CHARACTERISTICS:

LENGTH: 398 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single

TOPOLOGY: linear  
FEATURE:

NAME/KEY: CDS  
LOCATION: 1..398  
US-08-118-101A-5

Query Match: 7.3%  
Best Local Similarity: 63.6%  
Pred. No. 3, 4e-07;  
Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;

Score 53.2; DB 1;  
Length 398;  
Gaps 1;











[illegible]











[illegible]

PD	XX	26-OCT-2000.
XP	XX	14-APR-2000; 2000OWO-US10004.
PF	XX	15-APR-1999; 9905-0219404.
XX	XX	(UTAH ) UNIV UTAH RESS FOUN.
PA	XX	(UYVA ) UNIV YALE.
P1	XX	Abbott GW, Sealf F, Splawski I, Keating MT, Goldstein SAN;
P2	XX	WPI; 2000-617247/65.
DR	XX	P-PSDB; AAB29593.
PR	XX	Novel nucleic acids encoding MRP2, MRP3 and MRP4, useful for
PT	XX	diagnosing and treating ion channel disorders, especially long QT
PT	XX	syndrome -
PS	XX	Claim 56; Page 7; 132pp; English.
XX	XX	The invention relates to novel ion channel proteins related to
XX	XX	KCNH2, KIR2.1 and to nucleic acids encoding these proteins of
CC	XX	human and mouse KRCN2 (MRP2; AAB29587 and AAB29588,
CC	XX	respectively); human and mouse KRCN3 (MRP3; AAB29589 and AAB29588,
CC	XX	respectively). The cDNAs encoding these proteins are given in AAC64071-
CC	XX	AAC64076. KRCN2, along with HERG, forms cardiac fast delayed rectifier
CC	XX	potassium channels (I-IR), mutations in which are associated with long
CC	XX	QT syndrome. The invention also relates to methods of diagnosing long QT
CC	XX	syndrome using the KRCN2, KRCN3 or KRCN4 genes, a knockout mouse with a
CC	XX	defective KRCN2 gene, and a method of testing for long QT syndrome using
CC	XX	these proteins in animals comprising a heterologous ion channel protein gene
CC	XX	nothing animals comprising a heterologous ion channel protein gene
CC	XX	of the invention, a transgenic animal comprising human KRCN2 and HERG
CC	XX	DNA, and methods of and screening drugs for treating long QT syndrome
CC	XX	using KRCN2 proteins (including mutants), nucleic acids encoding them
CC	XX	and antibodies against KRCN2 proteins. The methods, antibodies, nucleic
CC	XX	acids, and proteins may be used for diagnosing or treating ion channel
CC	XX	disorders, especially long QT syndrome. Transgenic animals comprising
CC	XX	KRCN2 and HERG are useful for testing anti-long QT syndrome drugs.
CC	XX	KRCN2 and HERG are specifically claimed for use in diagnostic and drug screening
CC	XX	methods of the invention.
CC	XX	Note: The present sequence is not shown in the specification, but is
CC	XX	derived from the wild-type human KRCN2 cDNA sequence shown on page
XX	XX	118-119.
XX	XX	Sequence 732 BP; 221 A; 151 C; 156 G; 202 T; 0 other:
XX	XX	Query Match 99.8%; Score 730.4; DB 21; Length 732;
XX	XX	Best Local Similarity 99.94%; Pred. No. 9.2e-267;
XX	XX	Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
0y	1	CAAAATCCAGAAAGATGCGTTCCTTCACGTCGTCGCTATTTATTTAATTAATGCA 60
Db	1	CAAAATCCAGAAAGATGCGTTCCTTCACGTCGTCGCTATTTATTTAATTAATGCA 60
0y	61	GGAAGAGGAAACACATGCTACTTTATTCATTAATTCACAGACGCGGAAAGACGCTCCG 120
Db	61	GGAAGAGGAAACACATGCTACTTTATTCATTAATTCACAGACGCGGAAAGACGCTCCG 120
0y	121	AAGGATTTTATTTACTTATTAATGGAATTAATGAGCGCCGACAGACAAACAGCTACGAAAGGC 180
Db	121	AAGGATTTTATTTACTTATTAATGGAATTAATGAGCGCCGACAGACAAACAGCTACGAAAGGC 180
0y	181	CTCCGACACCCAAATGATGCTGCGAAGACTTCTATGATGATCTCTTACATGATGATGAT 240
Db	181	CTCCGACACCCAAATGATGCTGCGAAGACTTCTATGATGATCTCTTACATGATGATGAT 240
0y	241	GATTGGAATTTCTCTTTCATCATCTGGGACCTCGGAGACACGTGGAATTCGACCAAG 300
Db	241	GATTGGAATTTCTCTTTCATCATCTGGGACCTCGGAGACACGTGGAATTCGACCAAG 300

[illegible]

XX	RESULT 4
XX	ID AAC64084
XX	AAC64084 standard; DNA; 732 BP.
XX	AAC64084:
XX	19-FEB-2001 (first entry)
XX	Human potassium channel protein KCNE2 (MiRP1) M54T mutant DNA.
XX	Human: KCNE2; MiRP1, potassium channel protein; KCNE1-related;
XX	MiRP-related; long QT syndrome; cardiac arrhythmia;
XX	drug screening; knockout mouse; transgenic animal; ion channel
XX	fast delayed rectifier potassium channel; anti-KCNE2 antibody;
XX	NRK; mutant; ds.
XX	Homoduplex.
XX	Synthetic.
XX	NC200063434-AL.
XX	26-OCT-2000.
XX	14-APR-2000; 2000NC00510004.
XX	15-APR-1999; 99US-0128404.
XX	(UTAH ) UNIV UTAH RES FOUND.
XX	(UTAH ) UNIV YALE.
XX	Abbot GW, Sesti F, Splawski IJ, Keating MT, Goldstein SAN;
XX	WPI: 2000-67247/65.
XX	P-FSDB: AAC63594.
XX	Novel nucleic acids encoding MiRP1, MiRP2 and MiRP3, useful for
XX	diagnosing and treating ion channel disorders, especially Long QT
XX	syndrome -
XX	Claim 56: Page -/ 132pp; English.

XX The invention relates to novel ion channel proteins related to  
CC KCNE1 (M1NF) and to nucleic acids encoding them, the proteins of  
CC KCNE1 are human and rat KCNE2, hKCNK1, AAB25986, and AAB25986,  
CC KCNE2, AAB25987, and AAB25988,  
CC respectively), and human and mouse KCNE2 (M1NP3: AAB25989 and AAB25990,  
CC respectively), the cDNAs encoding these proteins are given in AAC4076-  
CC AAC4076, KCNE2, along with H1NF, forms cardiac fast delayed rectifier  
CC potassium channels (I<sub>Kr</sub>), mutations in which are associated with long  
CC QT syndrome. The invention also relates to methods of diagnosing with long  
CC QT syndrome.  
CC In the invention, a transgenic animal comprising human KCNE2 and HENG  
CC disruption in an endogenous KCNE2, KCNE2 or KCNE3 gene, transgenic  
CC nonhuman animals comprising a heterologous ion channel protein gene  
CC of the invention, a transgenic animal comprising human KCNE2 and HENG  
CC DNA, and methods of and secreting drugs for treating long QT syndrome  
CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
CC and antibodies against KCNE2 proteins, the methods, antibodies, nucleic  
CC acids and antibodies are used for diagnosing and treating long QT  
CC disorders, especially long QT syndrome, transgenic animals comprising  
CC KCNE2 and HENG are useful for testing anti-long QT syndrome drugs.  
CC The present sequence represents DNA encoding a mutant human KCNE2  
CC (M1NP3) specifically claimed for use in diagnostic and drug screening  
CC methods of the invention.  
CC The invention also relates to methods of diagnosing with long QT  
CC syndrome, the methods are not shown in the specification but is  
CC derived from the wild-type human KCNE2 cDNA sequence shown on page  
CC 118-119.

XX Sequence 732 BP; 221 A; 153 C; 157 G; 201 T; 0 other;

Query Match	Similarity	99.9%	Score	730.4	DB 21	Length	732
Best Local	Similarity	99.9%	Pred.	90.2e-203			
Matches	Conservative	1	Mismatches	1	Indels	0	Gaps
QY	1	CAATTCACGAAATATCCGTTTCTTCACATCTGTCGCTCC	DB	1	CAATTCACGAAATATCCGTTTCTTCACATCTGTCGCTCC	60	
QY	1	CAATTCACGAAATATCCGTTTCTTCACATCTGTCGCTCC	DB	1	CAATTCACGAAATATCCGTTTCTTCACATCTGTCGCTCC	60	
QY	121	AAGGATTTTATATCTATATATGAAATATGGGCGCGAAGACACACAGCGTACAGAGG	DB	61	CGAGAGGAGACATCTCATCTTATTCACACAGAGCGTACAGCGTCTCCG	120	
QY	121	AAGGATTTTATATCTATATATGAAATATGGGCGCGAAGACACACAGCGTACAGAGG	DB	61	CGAGAGGAGACATCTCATCTTATTCACACAGAGCGTACAGCGTCTCCG	120	
QY	121	AAGGATTTTATATCTATATATGAAATATGGGCGCGAAGACACACAGCGTACAGAGG	DB	121	AAGGATTTTATATCTATATATGAAATATGGGCGCGAAGACACACAGCGTACAGAGG	180	
QY	181	CGTCACGCGCAATGATGTACTGAAATCTCTCATATATGTCATCTCTACCTATGCTAT	DB	181	CGTCACGCGCAATGATGTACTGAAATCTCTCATATATGTCATCTCTACCTATGCTAT	240	
QY	241	GATTCGAAATCTCTCTTCTTCATCATCTGCGCCATCCTGTGACACATCTGCAAAAT	DB	241	GATTCGAAATCTCTCTTCTTCATCATCTGCGCCATCCTGTGACACATCTGCAAAAT	300	
QY	301	ACGGGAAACACTCAATGACCCCTGACACACCATATCAATTTGTAAGGACATGGACGAA	DB	301	ACGGGAAACACTCAATGACCCCTGACACACCATATCAATTTGTAAGGACATGGACGAA	360	
QY	361	CAGAGGCGCAATCTGATATATGAAATATGAAAGGCGCCACATCATGAAATATTTGG	DB	361	CAGAGGCGCAATCTGATATATGAAATATGAAAGGCGCCACATCATGAAATATTTGG	420	
QY	421	GCGTGGTTCAAAATGTCGCCCTGATATAGGGAAGAAAGGCAACCAAGTAACTGTGAC	DB	421	GCGTGGTTCAAAATGTCGCCCTGATATAGGGAAGAAAGGCAACCAAGTAACTGTGAC	480	
QY	481	CACATATGAAGATATCCATGCAATGCAAGCAATATGCTGCTGTATAGAGAA	DB	481	CACATATGAAGATATCCATGCAATGCAAGCAATATGCTGCTGTATAGAGAA	540	
QY	541	GTAATCTCTGCTCTGTTGTAGAAATTTCAATGAGCAATTAATGAGTGGCCATTAAGA	DB	541	GTAATCTCTGCTCTGTTGTAGAAATTTCAATGAGCAATTAATGAGTGGCCATTAAGA	600	
QY	541	GTAATCTCTGCTCTGTTGTAGAAATTTCAATGAGCAATTAATGAGTGGCCATTAAGA	DB	541	GTAATCTCTGCTCTGTTGTAGAAATTTCAATGAGCAATTAATGAGTGGCCATTAAGA	600	



XX Human potassium channel protein KCNE2 (MiRP1) R8A mutant DNA.  
 DE  
 XX  
 KW Human: KCNE2; potassium channel protein; KCNE1-related;  
 KW MiRP-related; long QT syndrome; cardiac arrhythmia;  
 KW drug screening; knockout mouse; transgenic animal; ion channel disorder;  
 KW fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
 KW HERG; mutant; DB.  
 OS  
 XX Homo sapiens.  
 XX Synthetic.  
 XX WO200063434-A1.  
 XX 26-OCT-2000.  
 XX 14-APR-2000; 2000MO-US10004.  
 XX 15-APR-1999; 99US-0129404.  
 XX (UTAH ) UNIV UTAH RES FOUND.  
 XX (UTAH ) UNIV YALE.  
 XX Abbott GW, Seethi F, Splawski I, Keating MT, Goldstein SAM;  
 DR WPI; 2000-672747/65.  
 DR P-PSDB; AAB293596.  
 XX Novel nucleic acids encoding MiRP1, MiRP2 and MiRP3, useful for  
 PF diagnosing and treating ion channel disorders, especially long QT  
 PT syndrome.  
 PS Claim 56: Page -: 132pp; English.

XX The invention relates to novel ion channel proteins related to  
 CC KCNE2 (MiRP) and to nucleic acids encoding them. The proteins of  
 CC this invention are human and rat KCNE2 (MiRP1, AAB293585 and AAB293586,  
 CC respectively), and human and mouse KCNE3 (MiRP2, AAB293587 and AAB293588,  
 CC respectively), and human and mouse KCNE4 (MiRP3, AAB293589 and AAB293590,  
 CC respectively). The cDNAs encoding these proteins are given in AAB24071-  
 CC AAB24076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
 CC potassium channels (I-KR), mutations in which are associated with long QT  
 CC syndrome. The invention also relates to methods of diagnosing long QT  
 CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
 CC nonhuman animal endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
 CC of the invention, a transgenic animal comprising human KCNE2 and human  
 CC DNA, and methods of and screening drugs for treating long QT syndrome  
 CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
 CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
 CC acids, and proteins may be used for diagnosing or treating ion channel  
 CC disorders, especially long QT syndrome. Transgenic animals comprising  
 CC the present invention are useful for testing anti-long QT syndrome drugs.  
 CC The present invention also relates to methods of diagnosing long QT  
 CC (MiRP1) specifically claimed for use in diagnostic and drug screening  
 CC methods of the invention.  
 CC Note: The present sequence is not shown in the specification, but is  
 CC derived from the wild-type human KCNE2 cDNA sequence shown on page  
 CC 118-119.  
 XX  
 XX  
 S0 Sequence 732 BP; 220 A; 152 C; 158 G; 202 T; 0 other:

Query Match 99.88; Score 730.4; DB 21; Length 732;  
 Beat Local Similarity 99.9%; Prod. No. 9.2e-203;  
 Matches 733; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

0Y 1 CAATTCAGAAAGATCGCTTTCTCACTTCGTGGCTATTTATTTAAATGCA 60  
 DB 1 CAATTCAGAAAGATCGCTTTCTCACTTCGTGGCTATTTATTTAAATGCA 60  
 61 GCAGAGGAGGACATGCTACTTATCAATTCACATTCACAAAGCGTGAAAGCTCTCCG 120  
 DB 61 GCAGAGGAGGACATGCTACTTATCAATTCACATTCACAAAGCGTGAAAGCTCTCCG 120

0Y 121 AAGCATTTTATTAATATATGAGCAATTGGCGCCGAGAACACACAGCTGAGCAAGGCC 180  
 DB 121 AAGCATTTTATTAATATATGAGCAATTGGCGCCGAGAACACACAGCTGAGCAAGGCC 180  
 0Y 181 CCGTCAGCAAGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 240  
 DB 181 CCGTCAGCAAGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 240  
 0Y 241 CCGTCAGCAAGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 300  
 DB 241 CCGTCAGCAAGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 300  
 0Y 301 ACGGGAAGACATCCATGACCCCTACACAGATGATGATGATGATGATGATGATGAT 360  
 DB 301 ACGGGAAGACATCCATGACCCCTACACAGATGATGATGATGATGATGATGATGAT 360  
 0Y 361 CAGACGCAATTCGATATGAGATGATGATGATGATGATGATGATGATGATGATGAT 420  
 DB 361 CAGACGCAATTCGATATGAGATGATGATGATGATGATGATGATGATGATGATGAT 420  
 0Y 421 GCGTGGGTTCAAAATGTCCTCCCTGATGATGATGATGATGATGATGATGATGATGAT 480  
 DB 421 GCGTGGGTTCAAAATGTCCTCCCTGATGATGATGATGATGATGATGATGATGATGAT 480  
 0Y 481 GCGTGGGTTCAAAATGTCCTCCCTGATGATGATGATGATGATGATGATGATGATGAT 540  
 DB 481 GCGTGGGTTCAAAATGTCCTCCCTGATGATGATGATGATGATGATGATGATGATGAT 540  
 0Y 541 GTGAGTCTCTGCTCTGTTGAGATGATGATGATGATGATGATGATGATGATGATGAT 600  
 DB 541 GTGAGTCTCTGCTCTGTTGAGATGATGATGATGATGATGATGATGATGATGATGAT 600  
 0Y 601 TGAATGACATTCATTCATGATGATGATGATGATGATGATGATGATGATGATGAT 660  
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 0Y 661 AAGCATTTTATTAATATATGAGCAATTGGCGCCGAGAACACACAGCTGAGCAAGGCC 720  
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 0Y 721 AAGCATTTTATTAATATATGAGCAATTGGCGCCGAGAACACACAGCTGAGCAAGGCC 780  
 DB 721 AAGCATTTTATTAATATATGAGCAATTGGCGCCGAGAACACACAGCTGAGCAAGGCC 780

RESULT 7  
 ID ABR6573 standard; DNA; 732 BP.  
 AC ABR6573;  
 XX  
 24-SEP-2002 (first entry)  
 DT  
 XX cDNA encoding human ether-a-go-go related interacting protein MiRP1.  
 KW Human: human ether-a-go-go related gene; HERG; KCR1; MiRP1;  
 KW long QT syndrome; LQT; single nucleotide polymorphism; cardiac arrhythmia;  
 KW potassium channel; ss; gene.  
 XX  
 XX Homo sapiens.  
 OS  
 XX Key Location/Qualifiers  
 XX CDS 7..443  
 FT /feature="a  
 FT /product="MiRP1"  
 PN WO200242735-A2.  
 XX  
 30-MAY-2002.  
 XX  
 30-OCT-2001; 2001MO-US45644.  
 PP





QY 121 AAGATTTTATTACTATATGAGCAATTGGGGGCGACAGCAACAGCTGAGGAGAGGC 180  
 DB 121 AAGATTTTATTACTATATGAGCAATTGGGGGCGACAGCAACAGCTGAGGAGAGGC 180  
 QY 181 AGGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 240  
 DB 181 AGGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 240  
 QY 241 GATTTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 300  
 DB 241 GATTTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 300  
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 DB 301 AGGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 360  
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 DB 361 CAGAGCCCAATCTTGATCTAGAGCAATGAGAGCCCAATCTAGAGCAATCTTGATCTAG 420  
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 DB 601 TTAGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 660  
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 DB 661 AAGACCTCTTACTCTCTGCGGCAAGTATGATGATGATGATGATGATGATGATGAT 720  
 QY 721 AAGACCAATTT 732  
 DB 721 AAGACCAATTT 732  
 RESULT 10  
 AAD35172 standard: DNA: 732 BP.  
 AAD35172:  
 25-JUL-2002 (first entry)  
 Human KCHN2 mutant DNA (T243C).  
 Human: Min-K related ion channel protein; MiRP1; ion channel disorder;  
 KCHN2; long QT syndrome; LQTS; cardiac arrhythmia; mutant; gene; SNP;  
 single nucleotide polymorphism; ds.  
 Homo sapiens.  
 Location/Dualifiers  
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 /tag="a"  
 /product="Human MiRP1 mutant protein"  
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 /standard\_name="Single nucleotide polymorphism (SNP)"  
 W0200222875-A2.  
 21-MAR-2002.

PF 11-SEP-2001; 2001MO-US28332.  
 PR 11-SEP-2000; 2000US-231571P.  
 XX (UYTA ) UNIV YALE.  
 XX Goldstein SAN;  
 PI MPI; 2002-362360/39.  
 DR P-PSDB; ABE22097.  
 XX Novel gene encoding Min-K related ion channel protein subunit and  
 PT polymorphisms in this gene associated with antiobesity-induced long QT  
 PS syndrome.  
 PS Claim 15; Page 46; 49pp; English.  
 CC The present invention relates to novel KCHN2 genes encoding Min-K related  
 CC (MiRP1) ion channel proteins and polymorphisms in these genes that  
 CC associated with ion channel disorders including antiobesity-induced long  
 CC QT syndrome, polymorphic ventricular tachycardia and other problems.  
 CC 57, 6116 of MiRP1 polymorphisms at nucleotide positions 11, 34,  
 CC encoding the amino acid positions is useful for diagnosing the presence  
 CC of a polymorphism that causes drug-induced LQTS. The diagnostic methods  
 CC are useful in the development of new drug therapies which selectively  
 CC target one or more KCHN2 polymorphisms that are associated with cardiac  
 CC arrhythmias. The present sequence is human KCHN2 mutant DNA (T243C).  
 XX  
 XX Sequence 732 BP; 221 A; 153 C; 157 G; 201 T; 0 other:  
 Query Match 99.8%; Score 730.4; DB 24; Length 732;  
 Best Local Similarity 99.9%; Pred. No. 9.2e-203;  
 Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 CAATTCGAAAGATCGTTTCTGTAACCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 60  
 DB 1 CAATTCGAAAGATCGTTTCTGTAACCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 60  
 QY 61 CAGAGAGGAGACATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 120  
 DB 61 CAGAGAGGAGACATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 120  
 QY 121 AAGATTTTATTACTATATGAGCAATTGGGGGCGACAGCAACAGCTGAGGAGAGGC 180  
 DB 121 AAGATTTTATTACTATATGAGCAATTGGGGGCGACAGCAACAGCTGAGGAGAGGC 180  
 QY 181 CCTTCACCAACCAATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 240  
 DB 181 CCTTCACCAACCAATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 240  
 QY 241 GATTTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 300  
 DB 241 GATTTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 300  
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 DB 301 ACGGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 360  
 QY 361 CAGAGCCCAATCTTGATCTAGAGCAATGAGAGCCCAATCTAGAGCAATCTTGATCTAG 420  
 DB 361 CAGAGCCCAATCTTGATCTAGAGCAATGAGAGCCCAATCTAGAGCAATCTTGATCTAG 420  
 QY 421 GGGTGGGTCTAAATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 480  
 DB 421 GGGTGGGTCTAAATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 480  
 QY 481 CAGACATGAGAGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 540  
 DB 481 CAGACATGAGAGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 540  
 QY 541 GTGAGTCTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCT 600  
 DB 541 GTGAGTCTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCT 600

07 601 TGAATGACATTCATCTAGATGATTTACTGCTGTGTGGAGCAAAATTTTGTCTGTA 660  
 Db 601 TGAATGACATTCATCTAGATGATTTACTGCTGTGTGGAGCAAAATTTTGTCTGTA 660  
 07 661 AGACCTCTTTTACTCTGGGCAAGATGTCATTTTAAATCAATCAATGATGAAT 720  
 Db 661 AGACCTCTTTTACTCTGGGCAAGATGTCATTTTAAATCAATCAATGATGAAT 720  
 07 721 AAAGCCAAATTT 732  
 Db 721 AAAGCCAAATTT 732  
 RESULT 11  
 ABO03173 standard: DNA: 732 BP.  
 AC ABO03173:  
 XX 25-JUL-2002 (first entry)  
 XX Human KCNE2 mutant DNA (A95G).  
 XX Human: Min-K related ion channel protein; MiRP1; Ion channel disorder;  
 KM KCNE2; long QT syndrome; LQTS; cardiac arrhythmia; mutant; gene; SNP;  
 KM single nucleotide polymorphism; ds.  
 XX Homo sapiens.  
 OS  
 XX Key  
 KM CDS  
 FT 732-815  
 FT /tag- a  
 FT /product- "Human MiRP1 mutant protein"  
 FT /replace (95, A)  
 FT /tag- b  
 FT /standard\_name- "Single nucleotide polymorphism (SNP)"  
 WT00022875-A2.  
 XX 21-MAR-2002.  
 PD 11-SEP-2001: 2001WO-US28332.  
 PE 11-SEP-2000: 2000US-231571P.  
 XX (UYVA ) UNIV YALE.  
 XX Goldstein SAN;  
 XX WPI: 2002-362360/39.  
 DR P-PSDB; ABE22098.  
 XX Novel gene encoding Min-K related ion channel protein subunit and  
 PT polymorphisms in this gene associated with antibiotic-induced long QT  
 syndrome  
 XX  
 XX Claim 18; Page 47-48; 49pp; English.  
 CC The present invention relates to novel KCNE2 genes encoding Min-K related  
 CC associated ion channel, proteins and polymorphisms in these genes that are  
 CC associated with long QT syndrome including antibiotic-induced long  
 CC QT syndrome (LQTS) detecting mutations at nucleotide position 44,  
 CC 57 or 116 of MiRP1 polypeptide or a mutation at a nucleotide position  
 CC encoding the amino acid positions is useful for diagnosing the presence  
 CC of a polymorphism that causes drug-induced LQTS. The diagnostic methods  
 CC are useful in the development of new drug therapies which selectively  
 CC target one or more KCNE2 polymorphisms that are associated with cardiac  
 CC arrhythmias. The present sequence is human KCNE2 mutant DNA (A95G).  
 XX  
 XX Sequence 732 BP: 220 A; 152 C; 158 G; 202 T; 0 other;  
 S0  
 Query Match 99.8%; Score 730.4; DB 24; Length 732;

Best Local Similarity 99.9%; Pred. No. 9,2e-203;  
 Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 07 1 CAAATCCAGAAAGAACCGCTTTCCTTAAACCTGTCGCTATTTATTTAATTTGCA 60  
 Db 1 CAAATCCAGAAAGAACCGCTTTCCTTAAACCTGTCGCTATTTATTTAATTTGCA 60  
 07 61 GCAAGAGGAGAACATGTCATCTTATTCATATTTCAACAAAGCGTGAGAACGCTTCG 120  
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 07 121 AAGGATTTTATTTACTTATGAGCAATTTGGCGCGAGAACACACAGCTGACAGAGG 180  
 Db 121 AAGGATTTTATTTACTTATGAGCAATTTGGCGCGAGAACACACAGCTGACAGAGG 180  
 07 181 CCTCCAAAGCCAAAGTGATGCTGTAAGAACTCTACTATGTCATCTGTCACCTAGTGAT 240  
 Db 181 CCTCCAAAGCCAAAGTGATGCTGTAAGAACTCTACTATGTCATCTGTCACCTAGTGAT 240  
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 Db 241 GATTGGATGTCCTCTTCATCATCTGTCGCACTCGGTGATCAGTGGAAATCCAAAG 300  
 07 301 AGCGGAACATCTCCAAATGACCCCTACACACATGATGTAAGAGATGCGAGGAAAGTA 360  
 Db 301 AGCGGAACATCTCCAAATGACCCCTACACACATGATGTAAGAGATGCGAGGAAAGTA 360  
 07 361 CAAAGCCAAATCTGATCTGATAGAAATGCAAGAGCCACATCATGATGAACATTTGTG 420  
 Db 361 CAAAGCCAAATCTGATCTGATAGAAATGCAAGAGCCACATCATGATGAACATTTGTG 420  
 07 421 GCGTGGTTCAAAATGTCCTCCCTGATGAGGAAAGAGCCACAGTACATCTGACATC 480  
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 07 541 GTGAGTCTGCTGCTCTTGTGTGAAATTTTGTATGAGATTTATGTTGGCCAAATAGA 600  
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 07 601 TGAATGACATTCATCTAGATGATTTACTGCTGTGTGGAGCAAAATTTTGTCTGTA 660  
 Db 601 TGAATGACATTCATCTAGATGATTTACTGCTGTGTGGAGCAAAATTTTGTCTGTA 660  
 07 661 AGACCTCTTTTACTCTGGGCAAGATGTCATTTTAAATCAATCAATGATGAAT 720  
 Db 661 AGACCTCTTTTACTCTGGGCAAGATGTCATTTTAAATCAATCAATGATGAAT 720  
 07 721 AAAGCCAAATTT 732  
 Db 721 AAAGCCAAATTT 732  
 RESULT 12  
 ABO03192 standard: cDNA: 600 BP.  
 AC ABO03192:  
 XX 11-JAN-2002 (first entry)  
 DE Human MiRP1 homologue-encoding cDNA, SEQ ID NO:968.  
 KM Human: cytokine; cell proliferation; cell differentiation; growth factor;  
 KM haematopoietic regulation; tissue growth; immunomodulator; actinin;  
 KM inhibin; chemokinesis; chemokinesis; thrombopoiesis; oncogenesis;  
 KM proliferation; metastasis; cancer; tumour; haematopoietic disorder;  
 KM myeloid cell disorder; lymphoid cell disorder; arthritis; arthritis;  
 KM myeloid cell disorder; lymphoid cell disorder; arthritis; arthritis;  
 KM atherosclerosis; coronary heart disease; arterial ischemia;  
 KM











GenCore version 5.1.4.P5.4578  
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OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 13:44:14 ; Search time 1593.65 seconds

(without alignments)  
13367.622 Million cell updates/sec

Title: US-09-550-163-1

Perfect score: 732  
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Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapept 1.0

Searched: 205640 seqs, 1455102878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database :

Listing first 45 summaries

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1: gb\_da:\*  
2: gb\_hlg:\*  
3: gb\_in:\*  
4: gb\_cm:\*  
5: gb\_ov:\*  
6: gb\_ov:\*  
7: gb\_ph:\*  
8: gb\_ph:\*  
9: gb\_pr:\*  
10: gb\_ro:\*  
11: gb\_ro:\*  
12: gb\_sy:\*  
13: gb\_sy:\*  
14: gb\_sy:\*  
15: gb\_sy:\*  
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17: em\_hum:\*  
18: em\_in:\*  
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41: em\_hlg\_hum:\*

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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2	732	100.0	732	6	AF071002
3	730.4	99.8	732	6	AX406939
4	730.4	99.8	732	6	AX406943
5	730.4	99.8	732	6	AX406945
6	730.4	99.8	732	6	AX406947
7	730.4	99.8	732	6	AX406949
8	730.4	99.8	732	6	AX406951
9	730.4	99.8	732	6	AX406953
10	730.4	99.8	732	6	AX406955
11	730.4	99.8	732	6	AX406957
12	730.4	99.8	732	6	AX406959
13	730.4	99.8	732	6	AX406961
14	730.4	99.8	732	6	AX406963
15	730.4	99.8	732	6	AX406965
16	730.4	99.8	732	6	AX406967
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23	730.4	99.8	732	6	AX406981
24	730.4	99.8	732	6	AX406983
25	730.4	99.8	732	6	AX406985
26	730.4	99.8	732	6	AX406987
27	730.4	99.8	732	6	AX406989
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29	730.4	99.8	732	6	AX406993
30	730.4	99.8	732	6	AX406995
31	730.4	99.8	732	6	AX406997
32	730.4	99.8	732	6	AX406999
33	730.4	99.8	732	6	AX407001
34	730.4	99.8	732	6	AX407003
35	730.4	99.8	732	6	AX407005
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37	730.4	99.8	732	6	AX407009
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41	730.4	99.8	732	6	AX407017
42	730.4	99.8	732	6	AX407019
43	730.4	99.8	732	6	AX407021
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#### ALIGNMENTS

Result No.	Score	Query Match	Length	DB ID	Description
1	732	100.0	732	6	AX406941
2	732	100.0	732	6	AF071002
3	730.4	99.8	732	6	AX406939
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9	730.4	99.8	732	6	AX406953
10	730.4	99.8	732	6	AX406955
11	730.4	99.8	732	6	AX406957
12	730.4	99.8	732	6	AX406959
13	730.4	99.8	732	6	AX406961
14	730.4	99.8	732	6	AX406963
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38	730.4	99.8	732	6	AX407011
39	730.4	99.8	732	6	AX407013
40	730.4	99.8	732	6	AX407015
41	730.4	99.8	732	6	AX407017
42	730.4	99.8	732	6	AX407019
43	730.4	99.8	732	6	AX407021
44	730.4	99.8	732	6	AX407023
45	730.4	99.8	732	6	AX407025

Pred. No. is the number of results predicted by chance to have a

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 BASE COUNT 221 a 152 c 157 g 202 t  
 ORIGIN

Query Match 100.0% Score 732; DB 6; Length 732;  
 Ref. Match 100.0% Pct. Ident. 100.0%  
 Mismatches 0; Indels 0; Gaps 0;  
 Matches 732; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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 DB 241 GATTGGAATGTTCTGTCATGTCGTGCGCATCTGCTGTGATGACCTGTGAAATTCAGAG 300  
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RESULT 2  
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 DEFINITION  
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 ACCESSION  
 VERSION  
 KEYWORDS  
 SOURCE  
 ORGANISM  
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 Balaictya; Primates; Catarrhini; Hominidae; Homo.  
 REFERENCE  
 AUTHORS  
 Abbott,C.W., Seest,F., Spilwack,I., Buck,M.E., Lehmann,M.H.,  
 Timothy,K.W., Keeling,M.T. and Goldstein,S.A.  
 MRP1 forms Ikr potassium channels with HERG and is associated with  
 cardiac arrhythmia  
 Cell 97 (2), 175-187 (1999)  
 JOURNAL  
 MEDLINE  
 99231923  
 12392319  
 REFERENCE  
 TITLE  
 AUTHORS  
 JOURNAL  
 Direct Submission  
 Submitted (05-JUN-1998) Section of Developmental Biology and  
 Biophysics, Department of Pediatrics and Boyer Center for Molecular  
 Medicine, Yale University School of Medicine, 295 Congress Avenue,  
 New Haven, Connecticut 06510  
 FEATURES  
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 /organism="Homo sapiens"  
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 /note="potassium channel subunit; MRP1"  
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 BASE COUNT 221 a 152 c 157 g 202 t  
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 Ref. Match 100.0% Pct. Ident. 100.0%  
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 DB 1 CAATCCGAAAGAAAGCCGTTTCCATACCTTGCGCATTTATTTAAATTTGA 60  
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 DB 181 CCGTCAGAGCAAGTGTATGAGACACTTCTACTGATCTGTCATGCTGATGCTGAT 240  
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 DEFINITION Sequence 1 from Patent WO222875.  
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 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE 1  
 AUTHORS Goldstein, S.A.  
 JOURNAL Polypeptide associated with cardiac arrhythmia  
 PUBLISHED 1992  
 PATENT NO 022875-A 1 21-MAR-2002;  
 YALE UNIVERSITY (US)  
 FEATURES  
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BASE COUNT 221 a 151 c 157 g 203 t  
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 Db 721 AAGGCCAAATTT 732

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 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.  
 REFERENCE 1  
 AUTHORS Goldstein, S.A.  
 JOURNAL Polypeptides associated with cardiac arrhythmia  
 PATENT: NO 022875-A 5 21-MAR-2002;  
 YALE UNIVERSITY (US)  
 FEATURES  
 source location/Qualifiers  
 1..732  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 74..445  
 /note="mink-related peptide 1, Thr substituted for Met at amino acid 54"  
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[illegible][illegible]





















(see [http://www.hgsc.bcm.tmc.edu/docs/genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html)).  
 NOTE: This is a 'working draft' sequence. It currently consists of 44 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as 'unknown' but the exact contig length is not known. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

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 1016 1115: gap of unknown length  
 1116 2238: contig of 1223 bp in length  
 2239 3486: gap of unknown length  
 3487 3786: gap of unknown length  
 3787 5609: contig of 1823 bp in length  
 5610 5709: gap of unknown length  
 5710 7138: contig of 1429 bp in length  
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Query Match 37.3%; Score 273; DB 2; Length 144709;  
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 Matches 329; Conservative 0; Mismatches 75; Indels 1; Gaps 1;

OY 56 CAGCAGAGGAGACATCTACTCTTTCATTCATTCATTCACAGACGCTGAGACAGCTTTC 118  
 Db 131731 CAGCAGAGGAGAGACATCTACTCTTTCATTCATTCATTCACAGACGCTGAGACAGCTTTC 131672

OY 119 CGAAGATTTTTATCTACTTATTTGAGACATTTGCCCCAGACACACAGCTGACAAAG 178  
 Db 131671 AAAAGGTTTATCTACTTATTTGAGACATTTGCCCCAGACACACAGCTGACAAAG 131612

OY 179 GCCTCCACAGCCAAATGATATGCTGAGACAACTTACTATGTCTACTGTACTGTATGCTG 238  
 Db 131611 GCCTCCACAGCCAAATGATATGCTGAGACAACTTACTATGTCTACTGTACTGTATGCTG 131552

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OY 419 CGCGAGGAGGTCGAAAATGTCGCGAGATGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 462  
 Db 131371 CGCGAGGAGGTCGAAAATGTCGCGAGATGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 131327

Search completed: May 21, 2003, 21:16:17  
 Job time : 2113.65 secs



GenCore version 5.1.4.P5.4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: May 15, 2003, 14:15:58 ; Search time 13 Seconds

(without alignments)  
392,430 Million cell updates/sec

Title: US-09-550-163-2

Perfect score: 632  
Sequence: 1 MSLTSMFTQCLEDFNRAIRIT.....EBSKATIHENICAGKFRKSP 123

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 112892 seqs, 41476328 residues

Total number of hits satisfying chosen parameters: 112892

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt\_40.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

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1	63.2	100.0	123	1 MRL_HUMAN	O9y616 homo sapien
2	55.4	87.7	123	1 MRL_RAT	G9w610 ratius norv
3	52.5	87.0	123	1 MRL_HUMAN	P15382 homo sapien
4	52.5	21.0	123	1 MRL_HUMAN	G9w610 ratius norv
5	52.5	20.3	129	1 MINK_MOUSE	P23450 melis silive
6	52.5	20.3	129	1 MINK_MOUSE	P15383 ratius norv
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33	52.5	20.3	130	1 MINK_MOUSE	G9w610 ratius norv

## ALIGNMENTS

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AC	O9y616	2000 (Ref. 39)			
BT	30-MAY-2000 (Ref. 39)				
DT	15-JUN-2002 (Ref. 41)				
DE	Minimum potassium ion channel-related peptide 1 (MIRP1) (Mink-related peptide 1).				
GN	KCNJ2.				
OS	Homo sapiens (Human)				
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;				
OX	Mammalia; Primates; Carnivora; Hominoidea; Homo.				
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RP	SEQUENCE FROM N.A., VARIANTS LQ76 E-9; T-54 AND T-57, AND VARIANT A-8.				
RC	TISSUE=Heart;				
RX	ABDOTT G.W., Sestl F., Splawski I., Buck M.E., Lejman M.H.,				
RA	McNally M., Mendez G.L., Goldstein S.A.H.,				
RT	Wulfeck B., et al. (1999) The human potassium channels with IRG and is associated with				
RL	cardiac arrhythmia. "				
12	SEQUENCE FROM N.A.,				
RA	Domenech A., Bellavilla X., de la Luna S.;				
RT	Cloning of human MIRP1 cDNA.				
RN	Submitted (SRR-2000) to the EMBL/GenBank/DBD databases.				
RX	ASSOCIATION WITH KCNJ2/KCNJ3, AND TISSUE SPECIFICITY.				
RA	MEDLINE=20487128; PubMed=11034315.				
RT	Tinel N., Diocot S., Lauritzen I., Barthelemy J., Lazdunski M.,				
RA	Bozsoeto M.;				
RT	"W-type KCNJ2-KCNJ3 potassium channels are modulated by the KCNE2				
RT	subunit. "				
RT	Submitted (SRR-2000) to the EMBL/GenBank/DBD databases.				
CC	- FUNCTION: AUXILIARY PROTEIN THAT CO-ASSEMBLE WITH A POTASSIUM				
CC	CHANNEL ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE				
CC	STABILITY OF THE MULTIMERIC COMPLEX. KCNE2 CO-ASSEMBLE WITH KCNJ2				
CC	(HERG) TO FORM THE CARDIAC POTASSIUM (IR) CHANNEL. MAY ALSO				
CC	ESSENTIALLY IDENTICAL PROPERTIES TO THE CHANNEL UNDERLYING THE				
CC	CONDUCTANCE WHICH PLAYS A CRITICAL ROLE IN DETERMINING THE				
CC	SUPRATHRESHOLD ELECTRICAL EXCITABILITY OF NEURONS. MAY ACT AS A				
CC	REGULATORY SUBUNIT OF THE KCNJ2/KCNJ3 M-TYPE CHANNEL.				
CC	- SUBUNIT: ASSOCIATES WITH KCNJ2. MAY ASSOCIATE WITH KCNJ2 AND/OR				
CC	KCNJ3.				
CC	- TISSUE SPECIFICITY: HIGHLY EXPRESSED IN BRAIN, HEART, SKELETAL				
CC	MUSCLE, PANCREAS, TESTIS, OVARY, UTERUS, AND SPERMATOCYTES.				
CC	SIGNIFICANT EXPRESSION IS FOUND IN LIVER, OVARY, TESTIS, PROSTATE,				
CC	SMALL INTESTINE, AND LEUKOCYTES. VERY LOW EXPRESSION, NEARLY				
CC	UNDETECTABLE, IN LUNG AND SPLEEN.				
CC	- DISEASE: DEFECTS IN KCNE2 ARE THE CAUSE OF LONG QT SYNDROME TYPE 6				
CC	(LQTS6); A FORM OF HEART DISEASE CHARACTERIZED BY CARDIAC				

CC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Rattus.
NCBI_TaxID=10116;	
RN	(1)
RS	SEQUENCE FROM N.A.*
RT	STRUCTURE OF THE KCH2C PROTEIN THAT CO-ASSEMBLES WITH A POTASSIUM CHANNEL ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE STABILITY OF THE MULTIMERIC COMPLEX. KCH2C CO-ASSEMBLES WITH KCNH2 (HERG) TO FORM THE CARDIAC POTASSIUM (IKr) CHANNEL.
RA	Abdotti G.W., Seel J.P., Slesewski I., Buck M.E., Lehmann M.H., Timothy K.W., Keating M.T., Goldstein S.A.N.;
RE	"MIRP forms IKr potassium channels with HERG and is associated with cardiac arrhythmias."
RL	Ceal 97-175-187(1999)
SC	-1-
CC	CHANNEL ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE STABILITY OF THE MULTIMERIC COMPLEX. KCH2C CO-ASSEMBLES WITH KCNH2 (HERG) TO FORM THE CARDIAC POTASSIUM (IKr) CHANNEL.
CC	-1-
CC	-1- SUBUNIT: ASSOCIATES WITH KCNH2.
CC	-1- SIMILARITY: BELONGS TO THE KCH2C FAMILY OF POTASSIUM CHANNELS.
CC	This Swiss-Prot entry is copyright 1997, it is produced through collaboration between the Swiss Institute of Bioinformatics and the EMBL outstations on the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (see http://www.isb-sib.ch/announce/ or send an email to license@isb-sib.ch).
CC	
DR	EMBL; AF071003; AAC20087.1;-
DR	InterPro: IPRO000369; ISR.Channel.
DR	Pfam: PF02046; ISR.Channel.1.
DR	PRINTS; PR00166; KCNCCHANNEL.
KW	Ionic channel; Transmembrane; Ion transport; Voltage-gated channel; Glycoprotein; Phosphorylation.
KV	
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FT	DOMAIN 2..70 CYTOSOLIC (POTENTIAL).
FT	DOMAIN 3..123 CYTOSOLIC (POTENTIAL).
FT	CARBOHYD 29 123 N-LINKED (GLCNAC.) (POTENTIAL).
FT	CARBOHYD 26 29 N-LINKED (GLCNAC...) (POTENTIAL).
FT	MOD_RES 71 71 PHOSPHORYLATION (BY PKC) (POTENTIAL).
FT	MOD_RES 74 74 PHOSPHORYLATION (BY PKC) (POTENTIAL).
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DB	1 MTFLANLTGTQLDEAFKFFETPTTMSRNRNTTAAAGLAARFDANERFYVILLVMWIOGF 60
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DB	61 AFIVALLVSVKSRKHESODPHYIEDMOQRYSOHLHDSMAITHBNLGATCF 120
OY	121 MSP 123
DB	121 VSP 123
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CD	P15382;
DY	01-APR-1990 (rel. 14, Created)
DY	01-APR-1990 (rel. 14, Last sequence update)
DY	01-JUN-2002 (rel. 41, Last annotation update)
CS	Channel voltage-gated potassium channel protein (Minimal potassium channel) (MIRK).
GN	KCNK1.
CS	Homo sapiens (Human).
CS	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
CS	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.



[illegible]

	CC	(VGLUT1) TO FORM THE SLOWLY ACTIVATED DELAYED RECTIFIER CARBONIC
	CC	POTASSIUM (IKS) CHANNEL. THE OUTWARD CURRENT REACHES ITS STEADY
	CC	STATE ONLY AFTER 50 SECONDS (BY SIMILARITY).
	CC	-1- SUBUNIT ASSOCIATES WITH KCNQ1 (BY SIMILARITY).
	CC	-1- SUBUNIT ASSOCIATES WITH KIR6.2 (BY SIMILARITY).
	CC	-1- PKC PHOSPHORYLATION INHIBITS THE POTASSIUM CURRENT (BY
	CC	SIMILARITY).
	CC	-1- SIMILARITY: BELONGS TO THE KCNE FAMILY OF POTASSIUM CHANNELS.
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	CC	or send an email to <a href="mailto:license@ebi.sdb.ch">license@ebi.sdb.ch</a> ).
	DR	EMBL; U63404; A041532.1; "
	DR	InterPro: IPR000369; ISR.Channel.
	DR	Plan: PFO2086; ISR.Channel: 1.
	DR	Notes: PROSITE: PS00750; Ion transport; Voltage-gated channel;
	KM	Tyrosine phosphorylation site. Ion transport; Voltage-gated channel;
	KM	Glycoprotein; Phosphorylation.
	FT	DOMAIN 1 43 EXTRACELLULAR (POTENTIAL).
	FT	TRANSMEM 44 66 POTENTIAL.
	FT	DOWNIN 67 129 CYTOPLASMIC (POTENTIAL).
	FT	CORNBARD 25 25 N-LINKED (GLYC.NC. ) (POTENTIAL).
	FT	CORNBARD 26 106 N-LINKED (GLYC.NC.BY RCS) (BY SIMILARITY).
	SO	SEQUENCE 129 AA; 14504 MW; F656A66CFDFDB95 CRC64;
		Query Match 20.6%; Score 130.5; DB 1; Length 129;
		Best Local Similarity 44.3%; Pred.No.2.9e-06;
		Matches 27; Conservative 15; Mismatches 16; Indels 3; Gaps 2;
	QY	51 LYXLMGMPSTIVVAITATKAKRENSDPDYDTED-NQRKR--QLNLESR 107
	Db	43 LYLWLVGFPGFPLGLMLSYRKRLKHSHDPDVEYSIQTKQKKAVLDANVLSYA 104
	QY	108 A 108
	QY	105 A 105
		RESULT 5
	MINI_MOUSE	
	ID	MINK_MOUSE STANDARD; PRF: 129 AA.
	AC	P23299;
	DT	01-NOV-1991 (Rel. 20, Created)
	DT	01-NOV-1991 (Rel. 20, Last sequence update)
	DT	30-MAY-2000 (Rel. 39, Last annotation update)
	DR	KM:slow voltage-gated potassium channel, protein (Minimal potassium
	RN	channel) (AIHK).
	RN	[1]
	OX	SEQUENCE FROM N.Y., AND PARTIAL SEQUENCE.
	OC	PTSDSPUR
	OC	RETRUSUR
	OC	MEDLINE:92007723; PubMed:1655403;
	RA	Honore E., Attali B., Romey G., Heurteaux C., Ricard P., Lesage F.,
	RA	Lazdunski M., Barhanin J.;
	RT	"Cloning, expression, pharmacology and regulation of a delayed
	RT	rectifier K <sup>+</sup> channel in mouse heart.";
	RL	EMBO J. 10:2805-2811(1991).
	RN	SEQUENCE FROM N.A.
	RN	RETRUSUR
	RN	MEDLINE:92233038; PubMed:1568475;
	RX	Lesage F., Lazdunski M., Barhanin J.;
	RX	ISR, a slowly activating voltage-sensitive K <sup>+</sup> channel.
	RT	Characterization of multiple cDNAs and gene organization in the



KM	Glycoprotein; 1	44	EXTRACELLULAR (POTENTIAL).
FN	DOMAIN	1	POTENTIAL.
FT	TRANSFERRIN	45	POTENTIAL.
FT	CARBOHYD	5	N-LINKED (GLYCINIC) (POTENTIAL).
FT	CARBOHYD	26	N-LINKED (GLYCINIC) (POTENTIAL).
FT	MOD. RES	103	PHOSPHORYLATION (BY PKC) (BY SIMILARITY).
SO	SEQUENCE	130 AA; 14671 MW; 7671EDED72492D C8G64;	
OY	Query Match	20.2%	Score 127.5; DB 1; Length 130;
Dy	Best Local Similarity	42.8%;	Fred. No. 5.9e-06;
OY	MATCHES 26;	Conservative 16;	Mismatches 10;
Dy	106 A 106		Indels 3; Gaps 2;
OY	106 A 106		
Dy	106 A 106		
RESULT 8			
MINK_CAVPO			
ID	MINK_CAVPO	STANDARD;	PRT; 125 AA.
AC	Q80A03; Q80A25; 1	39	(Created)
MC	30-MAY-2000 (Rel. 3);		Last sequence update)
DE	16-OCT-2001 (Rel. 40;		Last annotation update)
DE	ISK slow voltage-gated potassium channel protein (Musmal potassium channel) (MIM).		
GN	KCNK1.		
OS	Bavia porcellus (Guinea pig).		
OS	Eukaryota; Metazoa; Chordata; Carnivora; Vertebrata; Eutelestomii;		
OS	Phyla; Mammalia; Rodentia; Hypodigmahii; Cavidae; Cavia.		
OX	KCN1_Taxid=10141;		
OX	[1]		
RN	SEQUENCE FROM N.A.		
RP	TSSISE-Heart;		
RC	MEDLINE-94089666; PubMed-8265983;		
RA	Varnum M.D., Busch A.E., Bond C.T., Mayle J.J., Adelman J.P.;		
RA	"Fast activation of delayed rectifier K <sup>+</sup> current Iks and		
RL	mediates specific electrophysiological responses in adult guinea pigs"		
RL	Proc. Natl. Acad. Sci. U.S.A. 90:11528-11532(1993).		
RP	[2]		
RN	SEQUENCE FROM N.A.		
RP	TSSISE-Heart muscle;		
RC	MEDLINE-94173910; PubMed-7510407;		
RA	Zhang Z., Jurkiewicz N.K., Fohlman K., Lazarides E., Salata J.J.,		
RA	*alpha currents expressed from the guinea pig cardiac Isk protein are		
RL	enhanced by activators of protein Kinase C.*"		
RL	Proc. Natl. Acad. Sci. U.S.A. 91:1766-1770(1994).		
RT	-1- FUNCTION: ANGIOLATIN PROTEIN THAT CO-ASSEMBLES WITH A POTASSIUM		
CC	CHANNEL; ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE		
CC	(VOLTAGE) TO FORM THE SLOWLY ACTIVATING DELAYED RECTIFIER CARDIAC		
CC	ISK CURRENT IN GUINEA PIG HEART MYOCYTES BY RECRUITING ITS STANDY		
CC	STATE ONLY AFTER 50 SECONDS (BY SIMILARITY).		
CC	-1- SUBUNIT: ASSOCIATES WITH KCNQ1 (BY SIMILARITY).		
CC	-1- SUBCELLULAR LOCATION: Type I membrane protein.		
CC	-1- PTM: PHOSPHORYLATION INHIBITS THE POTASSIUM CURRENT (BY		
CC	SIMILARITY).		
CC	-1- SIMILARITY: BELONGS TO THE KCNc FAMILY OF POTASSIUM CHANNELS.		
CC	-----		
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CC	or send an email to licenses@ebi.ac.uk).		



D	DR	EMBL; L20462; AA972394.1;	-
OY		InterPro: IPR000369; ISK.Channel.	
DB		Pfam: PF02060; ISK.Channel; 1	
KM		Ionic channel; Transmembrane; Ion transport; Voltage-gated channel;	
RN		cytoplasmic phosphatidylserine phosphatase	
PT	PT	TRANSMEM	44 66 POTENTIAL
FT	FT	DOMAIN	67 125 CYTOPLASMIC (POTENTIAL) ..
FT	FT	CARBOHYD	5 5 N-LINKED (GLNACC..)
FT	FT	CARBOHYD	26 26 N-LINKED (GLNACC..)
FT	FT	CONFLICT	17 17 TYR -> SWY (IN REF. 1).
SQ		SEQUENCE	125 AA: ESGB84109ABSE76 CRG64
	Query Match	20.0%; Score 126.5; DB.1	Length 125;
	Best local similarity	44.9%; Pct-Id	7.1e-06;
	Matches	22; Conservative	15; Mismatches 11; Gaps 1;
RESULT 9			
ID	MINK_RABBIT	STANDARD:	PRT; 130 AA.
DT	30-MAY-2000 (rel. 39; Created)		
DI	30-MAY-2000 (rel. 39; Last sequence update)		
DE	30-MAY-2000 (rel. 39; Last annotation update)		
DF	ISK slow voltage-gated potassium channel protein (Minimal potassium channel) (MINK).		
GN	KCNEL.		
OS	Oryctolagus cuniculus (Rabbit).		
OC	Eukaryota; Metazoa; Chordata; Cenista; Vertebrata; Euteleostomi;		
OC	Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.		
RA	KCNL_Xtra019986;		
RP	SEQUENCE FROM N.A.		
RR	STRAIN-New Zealand white;		
RA	Fernald B., Jurkiewicz N.K., Jow B., Guinasso P., Foldander K., Swanson R., Salata J.U.;		
CC	Evidence of the slow component of the cardiac delayed rectifier K <sup>+</sup> current (IKs) in rabbit ventricular myocytes. J Biol Chem 274(4):2532-2538, 1999.		
CC	FUNCTION: ANIONIC PROTEIN THAT CO-ASSEMBLES WITH A POTASSIUM CHANNEL ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE STABILITY OF THE MULTIMERIC COMPLEX. KCNCL CO-ASSEMBLES WITH KCNOL (FVUQ1) TO FORM THE SLOWLY ACTIVATING DELAYED RECTIFIER CARDIAC POTASSIUM (IKS) CHANNEL. THE OUTWARD CURRENT REACHES ITS STEADY STATE ONLY AFTER 50 SECONDS (BY SIMILARITY).		
CC	- SUBUNIT ASSOCIATES WITH KCNOL (BY SIMILARITY).		
CC	- FUNCTIONAL DOMAIN:		
CC	- PKA PHOSPHORYLATION INHIBITS THE POTASSIUM CURRENT (BY SIMILARITY).		
CC	- SIMILARITY:		
CC	-1- SIMILARITY: BELONGS TO THE KCNE FAMILY OF POTASSIUM CHANNELS.		
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DR	EMBL; LA1659; AA93505.1;		
DR	InterPro: IPR000369; ISK.Channel.		
PFam	PF02060; ISK.Channel; 1		
CC	Ionic channel; Transmembrane; Ion transport; Voltage-gated channel;		
CC	cytoplasmic phosphatidylserine phosphatase		
FT	DOMAIN	1 44 EXTRACELLULAR (POTENTIAL) ..	
FT	DOMAIN	45 67 POTENTIAL	
FT	DOMAIN	68 130 CYTOPLASMIC (POTENTIAL).	

```

FT CARBOHYD 5 5 N-LINKED (GLCNAC, 1) (POTENTIAL)-
PT MOD_RES 103 103 PHOSPHORYLATION (BY PC) (BY SIMILARITY).
SQ SOURCE 130 AA: 14617 MW: 8710048742056 CRG6;

Query Match 19 98% Score 125, DB 1: Length 130;
Pairwise Similarity 34.24% PctId 95.16; Idents 12; Gaps 3;
Matches 25; Conservative

Db 106 GCCTVLENGLVAVH 118

0y 51 LTVAMVIGMSIVVAVSVMSRRSSNDYQVTV-VEPMQK-----YSSQILT-- 102
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 46 LTVAMVIGMSIVVAVSVMSRRSSNDYQVTV-VEPMQK-----YSSQILT-- 105
| : : : : : | : : : : : | : : : : : | : : : : : |

0y 103 ----IDSKATTH 111
| : : : : : | : : : : : | : : : : : | : : : : : |

Db 106 GCCTVLENGLVAVH 118

RESULT 10
ID Y204_METTA STANDARD: PRT: 439 AA.
Y204_METTA
1 204 100.00
2 100.00
3 100.00
4 100.00
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342 100.00
343 100.00
344 100.00
345 100.00
346 100
```



R		RP	SEQUENCE FROM N.A.	
RA	Nicholas J.;			
RL	Submitted (JAN-1996) to the EMBL/Genebank/DBJ databases.			
CC	-1- SIMILARITY : BELONGS TO FAMILY THAT GROUPS TOGETHER HSV-1 UL37,			
D	ENV-1 Z5; EBV BGLF1, VZV L1, HVS-1 G3, AND HCMV UL47.			
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KW	EMBL: U43400; AAC5692.1 ; -			
DR	Capid assembly.			
SQ	SEQUENCE 938 AA; 110170 MW; F4E39A2BFDD3BC9 CRC64;			
OY	Query Match			
Bt	Best Local Similarity 27.0%; Score: 74; DB 1; Length 938:			
Matches	30; Conservative 19; Mismatches 56; Indels 6; Gaps 4			
OY	3 TLSTPOTLEDVFRRFTITTDWNLSADGALADVANNFYLYTLWMIGMSF 62			
Db	773 TLFNFIIVSRISNTTTVDNLSDICSEHFNQLDPSFENLVYLDAFLKRTNFNF 831	:  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :		
OY	63 IVALIIVSYTKRKHREHSNPYHOI IVEDNGKRYSOLINI_EESKATTH 111			
Db	832 SLASDLAKKLAVKKO--DTIKMLNVODDVVLAKSHMLEFKOKPTLI 879			
RESULT 15				
ID	NID2M_CHOCRC STANDARD: PRT; 497 AA.			
AC	P48903;			
AF	01-FEB-1996 (Rel. 33 Created)			
DF	01-FEB-1996 (Rel. 33 Sequence updates)			
DT	15-JUL-1999 (Ref. 38, Last annotation update)			
DE	NADH-ubiquinone oxidoreductase chain 2 [EC 1.6.5.3].			
GN	ND2 OR ND2.			
OS	Chondrus crispus (Carrageen).			
OG	Eukaryota; Rhodophyta; Florideophyceae; Gigartinales; Gigartineaceae;			
NC	Eukaryotes.			
CC	Chondrus			
LN	GenBank:U43400;2765;			
PC	SEQUENCE FROM N.A.			
RP	TISSUE=Algae;			
RX	MEDLINE:95341661; PubMed-7616569;			
LA	Ledebnic C., Boyen C., Richard O., Bonnard G., Griememberger J.M.,			
RT	Kloareg B.;			
AB	"Complete sequence of the mitochondrial DNA of the rhodophyte			
CH	Chondrus crispus (Gigartinales)". Gene content and genome			
RI	J. Mol. Biol. 250:484-495(1995).			
CC	-1- CATALYTIC ACTIVITY : NADH + ubiquinone = NAD(+) + ubiquinol.			
CC	-1- SUBCELLULAR LOCATION : Integral membrane protein. Mitochondrial			
CC	inner membrane.			
CC	-1- SIMILARITY : BELONGS TO THE COMPLEX 1 SUBUNIT 2 FAMILY.			
CC				
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EMBL:	Z47547; CAAG8519.1 ; -			
Interpro:	IPIPO01750: Oxidored_1.			
Pfam:	PF00361: oxidored_qf_1.			
Oxidoreductase:	NAD; Ubiquinone; Mitochondrion; Transmembrane.			
SWISSPROT:	497 AA; DBOOEF38033FF40CA CRC64;			
SPONSEUR:	497 AA;			









	RL	DNA Res 8:205-213(2001).	1.	-
	DR	EMBL; AF003595; BAB75938.1.	1.	-
	DR	InterPro: IPR003593; ABC_Atpase.	1.	-
	DR	InterPro: IPR001140; AAA_Atptase.	1.	-
	DR	InterPro: IPR003595; CIMP_binding.	1.	-
	DR	InterPro: IPR001950; TIF_SUT1.	1.	-
	DR	Pfam: PF00604; ABC_membrane.1.	1.	-
	DR	Pfam: PF00005; ABC_tran.1.	1.	-
	DR	Pfam: PF00027; CIMP_Binding.1.	1.	-
	DR	ProDom: PD004006; Abc_Transport.1.	1.	-
	DR	SMART: SMO032; ANAP.1.	1.	-
	DR	SMART: SMO032; ANAP.1.	1.	-
	DR	PROSITE: PS00211; ABC_TRANSPORTER.1.	1.	-
	DR	PROSITE: PS01118; SUT1.1; UNKNOWN.1.	1.	-
	KW	ATP-binding; Complete proteome.		
	SQ	SEQUENCE 1003 AA: 11718 MW: 12606343.29066GRA CRC64;		
	Query Match	11.94; Score 75; DB 16; Length 1003;		
	Best Local Similarity	23.04; Pct Id 30.		
	Matches 21; Conservative 20; Mismatches 30; Indels 54; Gaps			
	QY	20 ITTDMNNTTAASGALAKKADKNVLYLVLWM-----IQNS--FIIVAI 67		
	DB	548 LKELEINIDQLTG--TALTGLDA---VESVYIITLVFYSMDLTGLDTGFPIFTTL 602		
	OY	66 ILTFVSKSR---REHNDPHQTYE-----DNGEQSLILNEE 105		
	DB	603 IASFVSRLAKSKENKSESTOSIVFMGIQTVAQNIELRSFSGEYARVY----- 657		
	QY	106 SKATHTENTGAAGFK 120		
	DB	658 -----AAQGR 662		
	RESULT 11			
	ID	069504 PRELMIMNFAY; PTR; 171 AA.		
	AC	069504;		
	DC	01-NOV-1996 (TEMBJREL_01, Created)		
	DE	01-NOV-1996 (TEMBJREL_01, Last sequence update)		
	DF	01-MAR-2003 (EMBL01, Last annotation update)		
	DI	U23 protein.		
	GN	U23.		
	OS	Human herpesvirus (Type 7 / strain J1) (HHV7).		
	OC	Viruses; dsDNA viruses, no RNA stage; Herpesviridae;		
	CC	Alphaherpesvirinae; Simplexvirus.		
	NCBI_TaxID=57278;			
	RN	SEQUENCE FROM N.A.		
	RA	Nicholas J.:		
	RC	Submitted (Jan-1996) to the EMBL/GenBank/DDBJ databases.		
	RN	[2]		
	RC	SEQUENCE FROM N.A.		
	RC	STRAIN-RK:		
	RA	Megaw A.G., Rapoport D., Avior B., Frenkel N., Davison A.J.:		
	RN	Submitted (Jan-1990) to the EMBL/GenBank/DDBJ databases.		
	RC	[3]		
	RN	SEQUENCE FROM N.A.		
	RC	STRAIN-RK:		
	RA	Megaw A.G., Frenkel N., Davison A.J.:		
	RN	Submitted (DEC-1997) to the EMBL/GenBank/DDBJ databases.		
	DR	EMBL; U43400; NC05684.1.		
	DR	EMBL; AF403218; AAC04736.1.		
	SQ	SEQUENCE 171 AA: 18663 MW: AD685FA509DDC20 CRC64;		
	Query Match	11.68; Score 73.5; DB 12; Length 171;		
	Best Local Similarity	36.68; Pct Id 6.3; Indels 3; Gaps		
	Matches 15; Conservative 11; Mismatches 3; Gaps 1.			
	50 IIYIIWWIGMGFT--IVAILVGVSKSRRESNDHYQY 87			







GenCore version 5.1.4.P5.4578  
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OM protein - protein search, using sw model

Run on: May 15, 2003, 14:25:44 ; Search time 53 Seconds  
(without alignments)  
223,839 Million cell updates/sec

Title: US-09-550-163-2

Sequence: 1 MSLSNFQTLQEDVFRRII.....ESKATIHENIGAGKRXSP 123

Scoring table: Gapop 10.0 , Gapext 0.5

Searched: 362588 segs, 96450795 residues

Total number of hits satisfying chosen parameters: 362588

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Maximum Match 0%

Listing first 43 summaries

Database :

Published Applications AA.\*  
1: /cgn2\_6/p/cdata2/pubpaa/US08\_NEM\_PUB pep.\*  
2: /cgn2\_6/p/cdata2/pubpaa/PCR\_NEM\_PUB pep.\*  
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14: /cgn2\_6/p/cdata2/pubpaa/US06\_PUBCOMB pep.\*  
Pred. No. is the number of results predicted by chance to have a score greater than or equal to the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Match	Length	DB ID	Description
1	632	100.0	123	9	US-10-000-151B-4
2	632	100.0	123	10	US-09-864-761-49007
3	533	84.3	103	10	US-09-864-761-36713
4	132.5	21.0	126	10	US-09-864-761-37234
5	132.5	21.0	126	10	US-09-815-242-12873
6	70	11.1	523	10	US-09-815-242-12723
7	70	11.1	525	10	US-09-815-242-12651
8	68.5	10.8	1590	9	US-10-180-326-1
9	67.5	10.7	380	10	US-09-134-333-12
10	67	10.6	272	10	US-09-815-242-5572
11	66	10.6	277	10	US-09-815-242-1282
12	65.5	10.4	553	10	US-09-815-242-1278
13	65.5	10.4	1276	10	US-09-882-610-24
14	65.5	10.4	2016	9	US-09-886-994-2
15	65.5	10.4	2016	10	US-09-840-125-4
16	65	10.3	367	10	US-09-815-242-10676
17	65	10.3	1482	10	US-09-815-242-12484
18	65	10.3	1835	10	US-09-935-541-5
19					

20	65	10.3	2175	10	US-09-935-541-2	Sequence 2, Appl1
21	65	10.2	2188	10	US-10-174-590-570	Sequence 4, Appl1
22	64.5	10.2	425	9	US-10-176-758-570	Sequence 570, App
23	64.5	10.2	425	9	US-10-176-758-570	Sequence 570, App
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45	64.5	10.2	425	9	US-10-175-737-570	Sequence 570, App

## ALIGNMENTS

RESULT 1  
US-10-000-151B-4  
; Sequence 4, Application US/10000151B  
; Publication No. US2003001316A1  
; GENERAL INFORMATION: Jeffrey R.  
; APPLICANT: George, Alfred L.  
; TITLE OF INVENTION: HUMAN KAI REGULATION OF HERG POTASSIUM CHANNEL BLOCK  
; CURRENT APPLICATION NUMBER: US/10/000.151B  
; CURRENT FILING DATE: 2000-10-30  
; NUMBER OF SEQ ID NOS: 5  
; SEQ ID NO 4  
; SEQ ID NO 4  
; LENGTH: 123  
; TYPE: PRT  
; ORGANISM: Homo sapiens  
US-10-000-151B-4  
Query Match 100.0% Score 632, DB 9, Length 123;  
Match 100.0% Positives 123, Gaps 0;  
Matches 123; Conservative 0; Mismatches 0; Indels 0;

QY 1 MSLSNFQTLQEDVFRRIIPTTMMNNRNTATACDAQKADENRYVYLIVLWIMCP 60  
DB I MSLSNFQTLQEDVFRRIIPTTMMNNRNTATACDAQKADENRYVYLIVLWIMCP 60  
QY 61 SFYVALIVSYVSKRHSNPNFHYIVEDQKTSVLSNLSKATIHENIGAGK 120  
DB 61 SFYVALIVSYVSKRHSNPNFHYIVEDQKTSVLSNLSKATIHENIGAGK 120  
QY 121 MSP 123  
DB 121 MSP 123  
QY III  
DB III  
RESULT 2  
US-09-864-761-49007  
; Sequence 49007, Application US/09864761  
; Patent No. US20020048763A1  
; GENERAL INFORMATION:  
; APPLICANT: Penn, Sharon G.



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1  OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 0.94
2  OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 0.59
3  OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 0.74
4  OTHER INFORMATION: EXPRESSED IN HEPA, SIGNAL = 0.66
5  OTHER INFORMATION: EXPRESSED IN SPLEEN, SIGNAL = 0.77
6  OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 0.66
7  OTHER INFORMATION: EST. HUMAN HIT: A1464239.1, EVALUE 2.00e-07
8  OTHER INFORMATION: SWISSPROT HIT: P15382, EVALUE 1.00e-39
9  US-09-864-761-37234
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11 Query Match 21.0% Score 132.5; DB 10; Length 76;
12 Best Local Similarity 42.1% Pred. No. 2.8e-07;
13 Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;
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15 Db 9 LVTAWMGESFFIVATLMSYKRSRRHSNDPFWYIESDANOEDKAVY 59
16
17 RESULT 5
18 US-10-338-316-4
19 Sequence 4 Application US/0138316
20 Publication No. US20020054380A1
21 GENERAL INFORMATION:
22 APPLICANT: Keating, Mark T.
23 APPLICANT: Sgambelli, Igor
24 APPLICANT: Sgambelli, Michael C.
25 TITLE OF INVENTION: MOTIONS IN THE KNEE! GENE ENCODING HUMAN MARK WHICH
26 TITLE OF INVENTION: KNEE! AS AN LQ7 GENE
27 TITLE OF INVENTION: KNEE! AS AN LQ7 GENE
28 FILE REFERENCE: 2233-162
29 CURRENT APPLICATION NUMBER: US/10/138, 316
30 CURRENT FILING DATE: 2002-05-06
31 PRIOR APPLICATION NUMBER: 09/444, 295
32 PRIOR FILING DATE: 1999-11-22
33 PRIOR APPLICATION NUMBER: 09/7135, 020
34 PRIOR FILING DATE: 1998-06-07
35 PRIOR APPLICATION NUMBER: 08/921, 068
36 PRIOR FILING DATE: 1997-08-29
37 PRIOR APPLICATION NUMBER: 08/739, 383
38 PRIOR FILING DATE: 1996-10-29
39 PRIOR APPLICATION NUMBER: 60/019, 014
40 PRIOR FILING DATE: 1995-12-22
41 PRIOR APPLICATION NUMBER: 09/094, 477
42 PRIOR FILING DATE: 1998-07-25
43 NUMBER OF SEQ ID NOS: 114
44 SOFTWARE: PatentIn Ver. 2.0
45 SEQ ID NO 4
46 LENGTH: 129
47 TYPE: PRT
48 ORGANISM: Homo sapiens
49 US-10-138-316-4
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51 Query Match 21.0% Score 132.5; DB 9; Length 129;
52 Best Local Similarity 45.1% Pred. No. 5.4e-07;
53 Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;
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55 Db 45 LVTAWMGESFFIVATLMSYKRSRRHSNDPFWYIESDANOEDKAVY 95
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57 RESULT 6
58 US-09-815-242-5723
59 Sequence 5723 Application US/09815242
60 Patent No. US20020061569A1
61 GENERAL INFORMATION:
62 APPLICANT: Hasebeek, Robert
63 APPLICANT: Kozlowski, John D.
64 APPLICANT: Zyskind, Judith W.
65 APPLICANT: Wall, Daniel
66 APPLICANT: Trawick, John D.
67 APPLICANT: Carr, Grant J.

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1 APPLICANT: Yamamoto, Robert T.
2 APPLICANT: Xu, H. Howard
3 TITLE OF INVENTION: Identification of Essential Genes in
4 THE INVENTION: Proteolyses
5 FILE REFERENCE: 1997-01-01
6 CURRENT APPLICATION NUMBER: US/09/815,242
7 PRIOR FILING DATE: 2001-03-21
8 PRIOR APPLICATION NUMBER: 66/191,076
9 PRIOR FILING DATE: 2000-03-21
10 PRIOR APPLICATION NUMBER: 66/206,848
11 PRIOR FILING DATE: 2000-05-23
12 PRIOR APPLICATION NUMBER: 66/207,727
13 PRIOR FILING DATE: 2000-05-26
14 PRIOR APPLICATION NUMBER: 66/215,442
15 PRIOR FILING DATE: 2000-10-23
16 PRIOR APPLICATION NUMBER: 66/251,625
17 PRIOR FILING DATE: 2000-11-27
18 PRIOR APPLICATION NUMBER: 66/257,931
19 PRIOR FILING DATE: 2000-12-22
20 PRIOR APPLICATION NUMBER: 66/269,308
21 PRIOR FILING DATE: 2001-02-16
22 NUMBER OF SEQ ID NOS: 14110
23 SOURCE: Genbank
24 SOFTWARE USED: For Windows Version 4.0
25 SEQ ID NO 5723
26 LENGTH: 523
27 TYPE: PR1
28 ORGANISM: Staphylococcus aureus
29 US-09-815-242-5723
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32 Best Local Similarity 11.1%; Score 70; DB 10; Length 523;
33 Matches 29; Conservative 14; Mismatches 33; Indels 28; Gaps
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RESULT 7  
US-09-815-242-12651  
Sequence 12651, Application US/09815242  
Patent No. US20020061569A1  
GENERAL INFORMATION:  
APPLICANT: YAMAMOTO, Robert  
APPLICANT: OHISEN, Karl E.  
APPLICANT: ZYKARD, Judith W.  
APPLICANT: WALL, Daniel  
APPLICANT: TRAWICK, John D.  
APPLICANT: CARR, Grant J.  
APPLICANT: YAMAMOTO, Robert T.  
APPLICANT: XU, H. Howard  
TITLE OR INVENTION: Identification of Essential Genes In  
Title OR INVENTION: Cytocycles  
FILE REFERENCE: ELIPPA 0114  
CURRENT FILING DATE: 2001-03-21  
PRIOR APPLICATION NUMBER: US/09/815,242  
CURRENT FILING DATE: 2001-03-21  
PRIOR APPLICATION NUMBER: 60/191,078  
PRIOR FILING DATE: 2000-03-21  
PRIOR APPLICATION NUMBER: 60/206,848  
PRIOR FILING DATE: 2000-05-23  
PRIOR APPLICATION NUMBER: 60/207,727  
PRIOR FILING DATE: 2000-05-26  
PRIOR APPLICATION NUMBER: 60/207,565  
PRIOR FILING DATE: 2000-10-23  
PRIOR APPLICATION NUMBER: 60/242,578  
PRIOR FILING DATE: 2000-11-27  
PRIOR APPLICATION NUMBER: 60/253,625  
PRIOR FILING DATE: 2000-12-22  
PRIOR APPLICATION NUMBER: 60/257,931  
PRIOR FILING DATE: 2000-12-22  
PRIOR APPLICATION NUMBER: 60/269,308

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? PRIOR FILING DATE: 2001-02-16
? NUMBER OF SEQ ID NOS: 14110
? SOFTWARE: FASTSEQ for Windows Version 4.0
? SEQ ID NO 14651
? LENGTH: 525
? ORGANISM: Staphylococcus aureus
US-09-815-242-14651

Query Match      11.1%; Score 70; DB 10; Length 525;
Best Local Similarity 27.9%; Pctd. No. 18;
Matches 29; Conservative 14; Mismatches 33; Indels 28; Gaps 5
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46 ELALIVALEA--G--G--G--G--G--G--G--G--G--G--G--G--G--G--G--
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94 -----EYKTSQIL--NLEF--SKATIHNGIAGKPKSP 123
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93 VEREDDENVAKYVQPLAHNRLEHNRKKNLEAVESGQIGDKNPS 136

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RESULT 8
US-10-180-326-1
Sequence 1, Application US/10190326
Publication No. US2003004366A1
GROSSER, STEPHEN W.
APPLICANT: Steino, Shunmu
APPLICANT: Shibusaki, Tadao
APPLICANT: Ozaki, No. US2003004966A1
TITLE OF INVENTION: Protein Rlm2
FILE REFERENCE: P21573
CURRENT APPLICATION NUMBER: US/10/180, 326
CURRENT FILING DATE: 2002-06-27
PRIOR APPLICATION NUMBER: JP 268372/99
CURRENT FILING DATE: 1999-10-08-160> 5
NUMBER OF SEQUENCES: 1
SOFTWARE: PatentIn version 3.0
SEQ ID NO 1
LENGTH: 1590
TYPE: PRT
ORGANISM: Mus musculus
US-10-180-326-1

Query Match      10.88; Score 68.5; DB 9; Length 1590;
Best Local Similarity 22.98; Freq. No. 1e+02; 57; Indexes 13; Gaps 3;
Matches 289; Conservative 179; Mismatches 57;
1 MSLSNFGFLDEDFRFFRFFTHDMRWKMTVALQGAALAKVADENRYVYLLVWIMGF 60
Db      26 WPLSHLE - EE - KRILLAWDQKREKREKQSVLKKEE-----KFAQPTQWF 72
QY      61 SFIVAILVSYVKSRRHSSNDPQHYQVTEQWQETYSQILNLSRSKATIHENNGAA 117
Db      73 PEGSITELVANNVLOPQOKQPKNEKEPQTKHDFEATKEDQVKKMGESQGGQEDRGDA 129

RESULT 9
US-09-134-333-12
Sequence 12, Application US/09134333
Patent No. US20020076403A1
GENERAL INFORMATION:
APPLICANT: LONGACRE-ANDRE, SHIRLEY
APPLICANT: ROTH, CHARLES
APPLICANT: MOTO, FARIIDMANO
APPLICANT: BARNWELL, JOHN
APPLICANT: ARNOLD, JAMES
TITLE OF INVENTION: RECOMBINANT PROTEIN CONTAINING A C-TERMINAL FRAGMENT OF
FILE REFERENCE: 0660-0135-OXCP
CURRENT APPLICATION NUMBER: US/09/134, 333
CURRENT FILING DATE: 1999-04-18
PRIOR APPLICATION NUMBER: PCT/FR97/00290

```

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? EARLIER FILING DATE: 1997-02-14
? EARLIER APPLICATION NUMBER: P966/01822
? EARLIER FILING DATE: 1996-02-14
? NUMBER OF SEQ ID NOS: 14
? SOFTWARE: Patentln Ver. 2.1
? SEQ ID NOS: 1 Patentln Ver. 2.1
? LENGTH: 380
? TYPE: PRT
? ORGANISM: Plasmodium vivax-like sp.
? FEATURE:
? OTHER INFORMATION: Amino Acids 1-140-REGION I
? FEATURE:
? OTHER INFORMATION: Amino Acids 141-178-REGION II
? FEATURE:
? OTHER INFORMATION: Amino Acids 179-283-REGION III
? OTHER INFORMATION: Amino Acids 284-380-REGION IV
? US-09-134-333-12

Query Match
Best Local Similarity 10.7%; Score 67.5; DB 10; Length 380;
Matches 31; Conservative 17; Mismatches 28; Indels 51; Gaps 6;

OY 28 QWTT-AACEALAKVDANENFYVLLVMWIGSFYVALVYVSKRREHN---81
DB 2 OVTGASAEAPETLVPAGISDHYVLPALQY-----KTKKQLEHNNFNT 52
OY 82 -----DPYQIVYDQKYSQILNESEKAT--110
DB 53 NTDPLDSNRKRRVLELYLNSDLPKXYSGETIYD--PK--LIDLEKKKLDS 107
OY 111 HENIGAA 117
DB 108 IYATIGAS 114

RESULT 10
US-09-815-242-5572
? Sequence 5572; Application US/09815242
? Patent No. US2002061569A1
? GENERAL INFORMATION:
? APPLICANT: Haselbeck, Robert
? APPLICANT: Ohlsen, Karl L.
? APPLICANT: Eyskind, Judith W.
? APPLICANT: Trawick, John D.
? APPLICANT: Carr, Grant J.
? APPLICANT: Yamamoto, Robert T.
? TITLE OF INVENTION: Identification of Essential Genes in
? TITLE OF INVENTION: Prokaryotes
? FILE REFERENCE: EPIYR 011
? CURRENT APPLICATION NUMBER: US/09/815,242
? PRIOR APPLICATION NUMBER: 60/191,078
? PRIOR FILING DATE: 2000-03-21
? PRIOR APPLICATION NUMBER: 60/206,848
? PRIOR FILING DATE: 2000-05-23
? PRIOR APPLICATION NUMBER: 60/207,727
? PRIOR FILING DATE: 2000-05-26
? PRIOR APPLICATION NUMBER: 60/242,578
? PRIOR FILING DATE: 2000-10-23
? PRIOR APPLICATION NUMBER: 60/253,625
? PRIOR FILING DATE: 2000-11-27
? PRIOR APPLICATION NUMBER: 60/257,931
? PRIOR FILING DATE: 2001-02-06
? PRIOR APPLICATION NUMBER: 60/269,308
? NUMBER OF SEQ ID NOS: 14110
? SOFTWARE: FastSeq for Windows Version 4.0
? SEQ ID NO 5572
? LENGTH: 272
? TYPE: PRT

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? ORGANISM: Staphylococcus aureus
? US-09-815-242-5572

Query Match
Best Local Similarity 10.6%; Score 67; DB 10; Length 272;
Matches 31; Conservative 21; Mismatches 44; Indels 54; Gaps 5;

OY 12 EDVFRIFITVNDNQWMTAFDQALQAVDANENFYVLLVMWIGSFYVALVST 71
DB 67 EDVFTSKLTFEDGCHENVAAYEASISDGLK--YLDLVWVGRTNE--AVMDT 119
OY 72 VKSKRREHND-----PRHOY-----87
DB 120 WKGMEDLYKNNVYANIGVSFPEHELEALLAQVSIKRYINOVETHTGKRLKLAQ 179
OY 88 -IIVEDQKYSQILNESEKATIHENG 115
DB 180 HYVESNPILMAQIILN-DETKIDIQEIG 208

RESULT 11
US-09-815-242-12282
? Sequence 12282; Application US/09815242
? Patent No. US2002061569A1
? GENERAL INFORMATION:
? APPLICANT: Haselbeck, Robert
? APPLICANT: Ohlsen, Karl L.
? APPLICANT: Eyskind, Judith W.
? APPLICANT: Trawick, John D.
? APPLICANT: Carr, Grant J.
? APPLICANT: Yamamoto, Robert T.
? TITLE OF INVENTION: Identification of Essential Genes in
? TITLE OF INVENTION: Prokaryotes
? FILE REFERENCE: EPIYR 011
? CURRENT APPLICATION NUMBER: US/09/815,242
? PRIOR APPLICATION NUMBER: 60/191,078
? PRIOR FILING DATE: 2000-03-21
? PRIOR APPLICATION NUMBER: 60/206,848
? PRIOR FILING DATE: 2000-05-23
? PRIOR APPLICATION NUMBER: 60/207,727
? PRIOR FILING DATE: 2000-05-26
? PRIOR APPLICATION NUMBER: 60/242,578
? PRIOR FILING DATE: 2000-10-23
? PRIOR APPLICATION NUMBER: 60/253,625
? PRIOR FILING DATE: 2000-11-27
? PRIOR APPLICATION NUMBER: 60/257,931
? PRIOR FILING DATE: 2000-12-22
? PRIOR APPLICATION NUMBER: 60/269,308
? NUMBER OF SEQ ID NOS: 14110
? SOFTWARE: FastSeq for Windows Version 4.0
? SEQ ID NO 12282
? LENGTH: 277
? TYPE: PRT
? ORGANISM: Staphylococcus aureus
? US-09-815-242-12282

Query Match
Best Local Similarity 10.6%; Score 67; DB 10; Length 277;
Matches 31; Conservative 21; Mismatches 44; Indels 54; Gaps 5;

OY 12 EDVFRIFITVNDNQWMTAFDQALQAVDANENFYVLLVMWIGSFYVALVST 71
DB 71 EDVFTSKLTFEDGCHENVAAYEASISDGLK--YLDLVWVGRTNE--AVMDT 123
OY 72 VKSKRREHND-----PRHOY-----87
DB 124 WKGMEDLYKNNVYANIGVSFPEHELEALLAQVSIKRYINOVETHTGKRLKLAQ 183
OY 88 -IIVEDQKYSQILNESEKATIHENG 115

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DB 164 HYMSSTPLNNMQLIN-DETRKDINQDLS 212

RESULT 12  
US-09-134-333-13

Sequence 13, Application US/09134333  
Patent No. US20020076403A1  
APPLICANT: JONASCH-ANDER, SHIRLEY  
APPLICANT: ROTH, CHARLES  
APPLICANT: NATA, FARIABAANO  
APPLICANT: BARNWELL, JOHN  
APPLICANT: MENDELIS, KAMINI  
TITLE OF INVENTION: RECOMBINANT PROTEIN CONTAINING A C-TERMINAL FRAGMENT OF  
FILIPIN REPEATS  
CURRENT FILING DATE: 1999-04-18  
EARLIER FILING DATE: 1997-02-14  
EARLIER APPLICATION NUMBER: PCT/FR97/00290  
EARLIER FILING DATE: 1996-02-14  
EARLIER SEQ ID NOS: 14  
SEQ ID NO 13  
LENGTH: 380  
TYPE: PRT  
ORGANISM: Plasmodium vivax-like sp.  
FEATURE:  
OTHER INFORMATION: Amino Acids 1-140-REGION I  
OTHER INFORMATION: Amino Acids 141-178-REGION II  
FEATURE:  
OTHER INFORMATION: Amino Acids 179-283-REGION III  
FEATURE:  
OTHER INFORMATION: Amino Acids 284-380-REGION IV  
US-09-134-333-13  
Query Match  
Best Local Similarity 10.5%; Score 66.5; DB 10; Length 380;  
Matches 32; Conservative 18; Mismatches 26; Indels 51; Gaps 7;  
DB 28 QNTT-AEENLQAVDAENFYVILYVAMVGFSEFYVALVYVKKSRHSHNS--- 81  
DB 2 QTTTGGSEHSEHFLVPAIGSDVYVYKFLKGT-----KTKGDLNNHNAFNT 52  
DB 82 -----DPYH-----GYVCEQMKYQSQIMNFSKATIT--- 110  
DB 53 NITDMLDSLRKKNYFLEVJMSDLNPFVSSSGEYIID---PYV--LIDLEKKKKLIGS 107  
DB 111 HENIGAA 117  
DB 108 YATIGAS 114  
RESULT 13  
US-09-815-242-5778  
Sequence 5778, Application US/09815242  
Patent No. US20020061569A1  
APPLICANT: INFORMATION:  
APPLICANT: OHLSEN, ROBERT  
APPLICANT: OHLSEN, KARI L.  
APPLICANT: ZYKLAND, JUDITH W.  
APPLICANT: WALL, DANIEL  
APPLICANT: TRAWICK, JOHN D.  
APPLICANT: CAIR, GRANT J.  
APPLICANT: YAMAMOTO, ROBERT T.  
TITLE OF INVENTION: METHOD OF IDENTIFICATION OF ESSENTIAL GENES IN  
FILE REFERENCE: ELITRA 011A  
CURRENT APPLICATION NUMBER: US/09/815.242

CURRENT FILING DATE: 2001-03-21  
PRIOR APPLICATION NUMBER: 60/191,078  
PRIOR FILING DATE: 2000-05-23  
PRIOR APPLICATION NUMBER: 60/206,848  
PRIOR FILING DATE: 2000-05-23  
PRIOR APPLICATION NUMBER: 60/207,177  
PRIOR FILING DATE: 2000-05-26  
PRIOR APPLICATION NUMBER: 60/242,578  
PRIOR FILING DATE: 2000-10-23  
PRIOR APPLICATION NUMBER: 60/191,253,625  
PRIOR FILING DATE: 2000-11-17  
PRIOR APPLICATION NUMBER: 60/257,391  
PRIOR FILING DATE: 2000-12-22  
PRIOR APPLICATION NUMBER: 60/269,308  
PRIOR FILING DATE: 2001-02-16  
NUMBER OF SEQ ID NOS: 14110  
SOFTWARE: STABLEST for Windows version 4.0  
SEQ ID NOS: 553  
LENGTH: 553  
TYPE: PRT  
ORGANISM: Staphylococcus aureus  
US-09-815-242-5778  
Query Match  
Best Local Similarity 10.4%; Score 65.5; DB 10; Length 553;  
Matches 26; Conservative 24; Mismatches 44; Indels 31; Gaps 5;  
DB 9 QTTEDVFRRI---FTYMDWQNTQVTSQALQAVDAENFYVILY----- 52  
DB 190 QDINDVIEAIKKNSEFVLLAGNRSSEETNAIKRLVETNLPVVEFGAGVSRLEN 249  
DB 53 -LAWNIGF-----SFITVALVSTYKSRHSHNSDPHYVCEQMKYQSQIMNFS 105  
DB 250 HFFCRVGLERNGVDELKMKDDVNTIT-----GVDT-ETLASMNNLELETOIINIDE 301  
DB 106 SKATI 110  
DB 302 VQAEI 306  
RESULT 14  
US-09-982-610-24  
Sequence 24, Application US/09982610  
Patent No. US20020146420A1  
GENERAL INFORMATION:  
APPLICANT: Genentech, Inc.  
Bennett, Brian D.  
Loo, James M.  
Lee, James M.  
Mathews, William  
Tsai, Siao Ping  
Wood, William I.  
TITLE OF INVENTION: PROTEIN TYROSINE KINASE AGONIST ANTIBODIES  
NUMBER OF SEQUENCES: 45  
COMPILED BY: Genentech, Inc.  
ADDRESS: 460 Point San Bruno Blvd  
CITY: South San Francisco  
STATE: California  
COUNTRY: USA  
ZIP: 94080  
COMPILED READABLE FORM: 1.44 Mb floppy disk  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: MMAPLAIN (Genentech)  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/982.610  
FILING DATE: 17-Oct-2001  
CLASSIFICATION: <UNKNOWN>  
PRIOR APPLICATION NUMBER: 08/444,668  
FILING DATE: 1996-MAY-23



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? APPLICATION NUMBER: 08/222616
? FILING DATE: 04-Apr-1994
? ATTORNEY/AGENT INFORMATION:
? NAME: Lee, Wendy M.
? ADDRESS: 40, 378
? TELEPHONE: 415/225-1994
? TELECOMMUNICATION INFORMATION:
? TELEPHONE: 415/225-1994
? TELEFAX: 415/952-9881
? INFORMATION FOR SEQ ID NO: 24:
? SEQUENCE CHARACTERISTICS:
? TYPE: amino acids
? TOPOLOGY: Linear
? SEQUENCE DESCRIPTION: SEQ ID NO: 24:
US-09-982-610-24

? QUERY MATCH
? Best Local Similarity: 10.4%; Score 65.5; DB 10; Length 1276;
? Matches 31; Conservative 13; Mismatches 37; Indels 29; Gaps 5;
OY 24 DNRRKMTACDEALQAKVDKENYVYLWVMIGMSFTYVALVTSYKRRG-HSD 82
DB 532 EGRN-----EQALLNGTAVVG---VVLVLV-----IYVAVLCKRSGREAEYS 576
OY 83 PEHQVY-----EMQEKYSQILMESSKATHTENICAGP 119
DB 577 KHQVYLHGCTVYIDPTEDPDMVREFAKIDVYKLVETVIGAF 626

RESULT 15
US-09-896-994-2
? Sequence 2, Application US/09896994
? Best Local Similarity: 20.3%; Pred. No. 2.9e+02;
? Matches 26; Conservative 13; Mismatches 37; Indels 31; Gaps 6;
GENERAL INFORMATION:
? APPLICANT: Ken Stokes
? TITLE OF INVENTION: SYSTEMS AND METHODS FOR ENHANCING CARDIAC
? CORRESPONDENCE ADDRESS: SIGNAL SENSING BY CARDIAC PACEMAKERS THROUGH GENETIC TREATY
? NUMBER OF SEQUENCES: 12
? STREET: One Liberty Place
? CITY: Philadelphia
? STATE: PA
? COUNTRY: U.S.A.
? ZIP: 19103
? COMPUTER READABLE FORM:
? MEDIUM TYPE: floppy disk
? OPERATING SYSTEM: PC-DOS/MS-DOS
? SOFTWARE: Nordperfect 6.1
? CURRENT APPLICATION DATA:
? FILING DATE: 02-Jul-2001
? CLASSIFICATION: <Unknown>
? PRIOR APPLICATION DATA:
? PUBLICATION NUMBER: 09/514,907
? FILING DATE: <Unknown>
? ATTORNEY/AGENT INFORMATION:
? NAME: Paul K. Legard
? REGISTRATION NUMBER: 38,534
? TELECOMMUNICATION INFORMATION:
? TELEPHONE: 212/5684300
? TELEFAX: 212/5684330
? INFORMATION FOR SEQ ID NO: 2:
? SEQUENCE CHARACTERISTICS:
? LENGTH: 2016 amino acids
? TYPE: amino acid
? STRANDEDNESS: single
? TOPOLOGY: unknown

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? SEQUENCE DESCRIPTION: SEQ ID NO: 2:
US-09-896-994-2

? QUERY MATCH
? Best Local Similarity: 10.4%; Score 65.5; DB 9; Length 2016;
? Matches 26; Conservative 13; Mismatches 43; Indels 31; Gaps 6;
OY 4 LSNFTQLEDPVRRPIRYDMNQNTTAEQALQAKVDKENYVYLWVMIGMSFT 63
DB 363 LALFRLTQDCWRELY-----QQTLS--AKIYIFPLVPLG--SPY 403
OY 64 IYVALVTSYKRRGSDPQVYEDQEKY-----SQILMESSKATHT--ENIGA 116
DB 404 LVLVLVYNAVAYED-----NQTATYERKAKRQEMAKKHEHLLITROVDYSR 458
OY 117 AGFRNSP 123
DB 459 SSIEMSP 465

Search completed: May 15, 2003, 14:35:09
Job time : 73 secs

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PF 30-JAN-2001; 2001MO-US00670.  
 XX  
 XX 04-FEB-2000; 2000US-0180312.  
 PR 26-MAY-2000; 2000US-0207456.  
 PR 30-JUN-2000; 2000US-0608409.  
 PR 03-AUG-2000; 2000US-0632366.  
 PR 21-SEP-2000; 2000US-0234487.  
 PR 27-SEP-2000; 2000US-0236359.  
 PR 04-OCT-2000; 2000GB-0024263.  
 PA (MOLE-) MOLECULAR DYNAMICS INC.  
 XX  
 XX Penn SG, Hanzel DK, Chen W, Rank DR;  
 XX WPI; 2001-488901/53.  
 DR  
 XX  
 XX Human genome-derived single exon nucleic acid probes useful for  
 PT analyzing gene expression in human cervical epithelial cells -  
 XX  
 XX Claim 27; SEQ ID NO 23453; 487bp; English.  
 XX  
 CC The present invention relates to human single exon nucleic acid probes  
 CC (SNP; see A110068-A128459). The present sequence is a peptide encoded  
 CC by one such probe. The SNPs are derived from human HeLa cells. The SNPs  
 CC can be used to produce a single exon microarray, which can be used for  
 CC measuring human gene expression in a sample derived from human cervical  
 CC epithelial cells. By measuring gene expression, the probes are therefore  
 CC useful in grading and/or staging of diseases of the cervix, notably  
 CC cervical cancer.  
 CC Note: The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at ftp.wipo.int/pub/published\_pat\_sequences.  
 XX  
 SQ Sequence 123 AA:  
 Query Match 100.08; Score 632; DB 22; Length 123;  
 Best Local Similarity 100.08; Pct. No. 1, 9e-67;  
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

PR 04-FEB-2000; 2000US-0180312.  
 PR 26-MAY-2000; 2000US-0207456.  
 PR 30-JUN-2000; 2000US-0608409.  
 PR 03-AUG-2000; 2000US-0632366.  
 PR 21-SEP-2000; 2000US-0234487.  
 PR 27-SEP-2000; 2000US-0236359.  
 PR 04-OCT-2000; 2000GB-0024263.  
 PA (MOLE-) MOLECULAR DYNAMICS INC.  
 XX  
 XX Penn SG, Hanzel DK, Chen W, Rank DR;  
 XX WPI; 2001-476286/51.  
 DR  
 XX  
 XX Novel single exon nucleic acid probe used to measuring gene expression  
 PT in a human breast -  
 XX  
 XX Claim 27; SEQ ID NO 14934; 322bp; English.  
 XX  
 CC The present invention relates to novel single exon nucleic acid probes  
 CC (see A110010-A110067). The present sequence is a peptide encoded by one  
 CC such probe. The SNPs are derived from human breast tissue. The SNPs  
 CC are derived from a human breast sample where the probe hybridises with strictly to a  
 CC nucleic acid expressed in the human breast. The probes are useful for  
 CC predicting, diagnosing, grading, staging, monitoring and prognosing  
 CC diseases of the human breast, particularly those diseases with polygenic,  
 CC aetiology. The diseases include: breast cancer, disorders of development,  
 CC inflammatory diseases of the breast, fibrocystic changes, proliferative  
 CC breast diseases and non-cancerous tumours did not form part of the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at ftp.wipo.int/pub/published\_pat\_sequences.  
 XX  
 SQ Sequence 123 AA:  
 Query Match 100.08; Score 632; DB 22; Length 123;  
 Best Local Similarity 100.08; Pct. No. 1, 9e-67;  
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MSTLSNFTQLEDFERRFTTNNRONTTAAQALAKVDANFYYVILTMWMIQNF 60  
 DB 1 MSTLSNFTQLEDFERRFTTNNRONTTAAQALAKVDANFYYVILTMWMIQNF 60  
 QY 61 SFYIALIVSYVSKRRSHSNDFPHQYIVEDQKRYXSQILNLSKATIHENIGAGFK 120  
 DB 61 SFYIALIVSYVSKRRSHSNDFPHQYIVEDQKRYXSQILNLSKATIHENIGAGFK 120  
 QY 121 MSP 123  
 DB 121 MSP 123  
 DE Peptide #4876 encoded by probe for measuring breast gene expression.  
 XX  
 XX Probe: human; breast disease; breast cancer; development disorder;  
 KW inflammatory disease; proliferative breast disease; non-carcinoma tumour.  
 XX  
 XX Homo sapiens.  
 XX  
 XX WO200157270-A2.  
 XX  
 XX 09-AUG-2001.  
 XX  
 XX 29-JAN-2001; 2001MO-US00661.  
 XX

QY 1 MSTLSNFTQLEDFERRFTTNNRONTTAAQALAKVDANFYYVILTMWMIQNF 60  
 DB 1 MSTLSNFTQLEDFERRFTTNNRONTTAAQALAKVDANFYYVILTMWMIQNF 60  
 QY 61 SFYIALIVSYVSKRRSHSNDFPHQYIVEDQKRYXSQILNLSKATIHENIGAGFK 120  
 DB 61 SFYIALIVSYVSKRRSHSNDFPHQYIVEDQKRYXSQILNLSKATIHENIGAGFK 120  
 QY 121 MSP 123  
 DB 121 MSP 123  
 DE Amino acid sequence of human potassium channel subunit ISK2.  
 XX  
 XX Human; potassium channel; ISK2; gene therapy; gastric motility;  
 KW gastric acid secretion; anti-arrhythmic agent; myocardial infarction.  
 XX  
 XX Homo sapiens.  
 XX  
 XX WO200127246-A1.  
 XX  
 XX 19-APR-2001.  
 XX  
 XX 10-OCT-2000; 2000MO-US28014.  
 XX

ER 12-OCT-1999; 99US-0158781.  
 XX (MERK ) MERCK & CO INC.  
 XX Swanson RJ, Liu Y, Folander K;  
 PI WPI: 2001-273764/28.  
 DR N-PSDB; AAF80269.  
 XX New DNA encoding the 18k2 potassium channel subunit, useful e.g. for  
 PT detecting mutations and screening for therapeutic agents  
 PS Claim 8; Fig 1b; 46pp; English.  
 XX The present sequence represents a human potassium channel subunit,  
 CC designated 18k2. The 18k2 polynucleotide, and derived probes, are  
 CC specifically selected mutations in the 18k2 gene, to determine  
 CC levels of mRNA expression and to screen for mutations in the 18k2  
 CC recombinant expression of 18k2; in gene therapy to increase potassium  
 CC channel activity and to generate transgenic animals, as models for  
 CC for drug screening. Recombinant 18k2 is used for studying biochemical  
 CC activity of 18k2 and its role in disorders of gastric motility and  
 CC gastric acid secretion, and to raise specific antibodies. 18k2  
 CC is a eukaryotic protein useful for treating diseases associated with  
 CC increased gastric acid secretion, e.g. reflux, gastritis, and  
 CC anti-arrhythmic agents for treating myocardial infarction and as  
 CC regulators of gastric acid secretion.  
 XX Sequence 123 AA:  
 SQ  
 Query Match 100.0%; Score 632; DB 22; Length 123;  
 D Match Similarity 100.0%; Pred. No. 1,9e-67;  
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Oy 1 NKTLSNFTQTLDEYFRIFITYDMNMONTATADGALQAVDAENFYVILVLMYMGF 60  
 Db 1 NKTLSNFTQTLDEYFRIFITYDMNMONTATADGALQAVDAENFYVILVLMYMGF 60  
 Oy 61 SFTIVALLSTYKSRREHSNDYHOYIVDMQEKYSQILNLESKATHTENIGAGKR 120  
 Db 61 SFTIVALLSTYKSRREHSNDYHOYIVDMQEKYSQILNLESKATHTENIGAGKR 120  
 Oy 121 MSP 123  
 Db 121 MSP 123  
 RESULT 6  
 ID AAV00215 standard; Protein: 123 AA.  
 AC AAV00215;  
 DM 10-MAY-2001 (first entry)  
 XX Human potassium channel regulatory protein, MINK2.  
 XX Human; MINK2; potassium channel; cardiac arrhythmia; hypertension;  
 KW angina; asthma; diabetes; renal insufficiency; urinary incontinence;  
 KM irritable colon; epilepsy; cerebrovascular ischemia; autoimmune disease.  
 OS Homo sapiens.  
 XX Homo sapiens.  
 PN M0200114403-A1.  
 XX 01-MAR-2001.  
 PD 18-AUG-2000; 2000MO-0522799.  
 XX 20-AUG-1999; 99US-0378201.  
 PA (UYCA-) UNIV CASE WESTERN RESERVE.  
 XX

PI Flicker E, Wible B, Brown AM;  
 XX WPI: 2001-218424/22.  
 DR N-PSDB; AAS00245.  
 XX Novel potassium channel gene termed Mink2 encoding potassium channel  
 PT regulatory protein, useful for screening compounds that are useful for  
 PT treating diseases caused by aberrant potassium activity  
 XX Disclousure; Fig 9; 39pp; English.  
 PS  
 XX The sequence represents the amino acid sequence of human potassium  
 CC channel regulatory protein, MINK2. MINK2 sequence is useful for producing  
 CC a potassium channel regulatory protein useful for in vitro or in vivo  
 CC screening of agonistic or antagonistic compounds that are useful for  
 CC treating diseases caused by aberrant potassium activity, such as human  
 CC cardiac arrhythmia, hypertension, angina, asthma, diabetes, renal  
 CC insufficiency, urinary incontinence, irritable colon, epilepsy,  
 CC cerebrovascular ischemia, and autoimmune disease.  
 XX Sequence 123 AA:  
 SQ  
 Query Match 100.0%; Score 632; DB 22; Length 123;  
 D Match Similarity 100.0%; Pred. No. 1,9e-67;  
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Oy 1 NKTLSNFTQTLDEYFRIFITYDMNMONTATADGALQAVDAENFYVILVLMYMGF 60  
 Db 1 NKTLSNFTQTLDEYFRIFITYDMNMONTATADGALQAVDAENFYVILVLMYMGF 60  
 Oy 61 SFTIVALLSTYKSRREHSNDYHOYIVDMQEKYSQILNLESKATHTENIGAGKR 120  
 Db 61 SFTIVALLSTYKSRREHSNDYHOYIVDMQEKYSQILNLESKATHTENIGAGKR 120  
 Oy 121 MSP 123  
 Db 121 MSP 123  
 RESULT 7  
 ID AAV99168 standard; Protein: 123 AA.  
 AC AAV99168;  
 DM 24-SEP-2002 (first entry)  
 XX human ether-a-go-go related interacting protein MMRP1.  
 XX Human; human ether-a-go-go related gene; HERG; KCRI; MMRP1;  
 KM long QT syndrome; LQT; single nucleotide polymorphism; cardiac arrhythmia;  
 KW potassium channel.  
 OS Homo sapiens.  
 XX Homo sapiens.  
 PN M0200242735-A2.  
 XX 30-MAY-2002.  
 PD 30-OCT-2001; 2001MO-0545644.  
 XX 30-OCT-2000; 2000US-244340P.  
 XX (UYVA-) UNIV VANDERBILT.  
 XX Balcer JR, George AL, Roden DW;  
 PI WPI: 2002-527650/56.  
 DR N-PSDB; ABR86573.  
 XX Identifying a potassium channel activity modulator for drug design,  
 PT comprising a potassium channel and a potassium channel and creat  
 PT cerebral DNA library (KCRI) polypeptide, and determining activity -

XX Disclosure: Page 163; 164pp; English.

CC The invention relates to identifying (M1) a compound that modulates  
CC biological activity of a potassium channel (PC), by contacting a  
CC compound with a structure comprising a PC polypeptide and a polypeptide  
CC encoding a KChIP1 protein, in the presence of a KChIP1 protein, and  
CC measuring the activity of the PC polypeptide in the presence of the  
CC compound, where a difference in the activities indicates modulation of  
CC biological activity of PC. Also include are identifying (M2) a candidate  
CC compound that modulates the biological activity of a complex comprising a  
CC human ether-a-go-go-related gene (HERG) channel polypeptide and a KChIP1  
CC polypeptide, identifying (M3) a candidate compound as a modulator of KChIP1  
CC expression, modulating (M4) PC function in a subject, comprising  
CC administering to a subject a compound that modulates the activity of a  
CC KChIP1-encoding nucleic acid molecule in a cell or tissue where modulated  
CC PC function is desired, screening (M5) for susceptibility to a drug  
CC induced cardiac arrhythmia in a subject, comprising obtaining a  
CC biological sample from the subject and detecting a polymorphism of a KChIP1  
CC gene in the biological sample from the subject, where the presence of the  
CC polymorphism indicates the susceptibility of the subject to a  
CC drug-induced cardiac arrhythmia, identifying (M6) a first  
CC oligonucleotide of the pair hybridizes to a first portion of a KChIP1 gene  
CC which includes a polymorphism of the KChIP1 gene, and the second  
CC oligonucleotide of the pair hybridizes to a second portion of the KChIP1  
CC gene that is adjacent to the first portion and a set of antisense  
CC oligonucleotide primers, suitable for amplifying a portion of a KChIP1 gene  
CC which includes a polymorphism of the KChIP1 gene, (M1) is useful for  
CC identifying a compound that modulates the activity of a  
CC KChIP1-encoding nucleic acid molecule in a cell or tissue where modulated  
CC activity) in a mammal, by preparing a composition comprising the  
CC compound and administering the composition. The compound is useful for  
CC treating or preventing long QT syndrome (LQTS) and is useful in drug  
CC protein M1RP1 (not defined).

XX Sequence 123 AA:

Query Match 100.0%; Score 632; DB 23; Length 123;  
Best Local Similarity 100.0%; Pred. No. 1.9e-67;

Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MSTLSNFTQLEDFRFRITFYDMNQNTTAAQNALQKADANFYVILTLWMIQNF 60  
DB 1 MSTLSNFTQLEDFRFRITFYDMNQNTTAAQNALQKADANFYVILTLWMIQNF 60

QY 61 SFITVALIVSVTKSKRRHSNDPHYQYIVEDMQEKYSQILNIESKATIHENIGAAGFK 120  
DB 61 SFITVALIVSVTKSKRRHSNDPHYQYIVEDMQEKYSQILNIESKATIHENIGAAGFK 120

QY 121 MSP 123  
DB 121 MSP 123

XX 21-MAR-2002.

RESULT 8

AAE22095 AAE22095 standard; Protein; 123 AA.

AC AAE22095;

XX 25-JUL-2002 (first entry)

DE Human M1RP1 wild type protein.

XX Human; Min-K related ion channel protein; M1RP1; ion channel disorder;  
XX KChIP2; long QT syndrome; LQTS; cardiac arrhythmia.

OS Homo sapiens.

XX WO200222875-A2.

PN 21-MAR-2002.

XX 11-SEP-2001; 2001MO-US28332.

XX 11-SEP-2000; 2000US-231571P.

XX (UYUA ) UNIV YALE.

XX Goldstein SAN;

XX WPI; 2002-362360/39.

XX DR N-PSDB; AAD35170.

XX Novel gene encoding Min-K related ion channel protein subunit and  
XX polymorphisms in this gene associated with antiarrhythmic-induced long QT  
XX syndrome.

XX Claim 8; Fig 1A; 49pp; English.

CC The present invention relates to novel KChIP2 genes encoding Min-K related  
CC (M1RP) 1 ion channel proteins and polymorphisms in these genes that are  
CC associated with ion channel disorders including arrhythmic-induced long  
CC QT syndrome. The invention includes a first oligonucleotide of the pair  
CC 5' or 116 of M1RP1 polypeptide or a mutation at a nucleotide position  
CC encoding the amino acid position is useful for diagnosing the presence  
CC of a polymorphism that causes drug-induced LQTS. The diagnostic methods  
CC are useful in the development of new drug therapies which selectively  
CC target one or more KChIP2 polymorphisms that are associated with cardiac  
CC arrhythmias. The present sequence is human M1RP1 wild type protein.

XX Sequence 123 AA:

Query Match 100.0%; Score 632; DB 23; Length 123;  
Best Local Similarity 100.0%; Pred. No. 1.9e-67;

Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MSTLSNFTQLEDFRFRITFYDMNQNTTAAQNALQKADANFYVILTLWMIQNF 60  
DB 1 MSTLSNFTQLEDFRFRITFYDMNQNTTAAQNALQKADANFYVILTLWMIQNF 60

QY 61 SFITVALIVSVTKSKRRHSNDPHYQYIVEDMQEKYSQILNIESKATIHENIGAAGFK 120  
DB 61 SFITVALIVSVTKSKRRHSNDPHYQYIVEDMQEKYSQILNIESKATIHENIGAAGFK 120

QY 121 MSP 123  
DB 121 MSP 123

RESULT 9

ABBI1948 ABBI1948 standard; peptide; 135 AA.

XX ABBI1948;

XX 11-JAN-2002 (first entry)

DE Human M1RP1 homologue, SEQ ID NO:2318.

XX Human; cytokine; cell proliferation; cell differentiation; growth factor;  
XX haematopoietic regulation; tissue growth; immunomodulation; activin;  
XX inhibin; chemotaxis; chemokinesis; thrombolysis; oncogenesis;  
XX myeloid cell disorder; lymphoid cell disorder; asthma; arthritis;  
XX chronic inflammatory condition; proliferative retinopathy;  
XX atherosclerosis; coronary heart disease; arterial ischemia;

XX bone disorder; osteoporosis; vascular growth disorder; disorders;  
XX cell culture; drug screening; gene therapy; antiinflammatory;  
XX cell culture; drug screening; gene therapy; antiinflammatory;  
XX cytoskeletal; antiarrhythmic; haematostatic; antiarrhythmic;  
XX antifungal; vasodilatory; cardiatic; vitinoid; antibacterial;

XX Homo sapiens.





Oy	1	N6S8NFQTLDEYFRATITTYMDNNRONTFADLALAKDAENFYVYLTLAWIDHM	60
Db	61	SPTTIALIVSTYKSRHNSDTPQTGYVEDMGRYKSQLIMESKATHENTGACGF	120
Oy	61	SPTTIALIVSTYKSRHNSDTPQTGYVEDMGRYKSQLIMESKATHENTGACGF	120
Db	73	SPTTIALIVSTYKSRHNSDTPQTGYVEDMGRYKSQLIMESKATHENTGACGF	132
Oy	121	MSP 123	
Db	133	MSP 135	
ID	AAB29593	standard; Protein: 123 AA.	
AC	AAB29593;		
DT	19-FEB-2001	(first entry)	
DD	Human potassium channel protein KCNE2 (MiRP1) mutant Q9E.		
KM	Human: KCNE2; MiRP1; potassium channel protein; KNCB2-related;		
KM	Mink-related; long QT syndrome; cardiac arrhythmias;		
KM	drug screening; knockout mouse; transgenic animal; ion channel		
KM	fast delayed rectifier potassium channel; anti-KCNE2 antibody;		
KX	HERG; mutant; multiin.		
KX	hemolysates.		
OS	synthetic.		
PN	M0200063434-A1.		
PD	26-OCT-2000.		
PF	14-APR-1990; 2000MO-US10004.		
PE	15-APR-1999; 990US-0129404.		
PA	(UTAH) UNIV UTAH RES FOUND.		
XX	(UYVA) UNIV YALE.		
P1	Abdott GW, Seethi F, Splawski I, Keating MT, Goldstein SAN;		
DR	WPI: 2000-672747/65.		
XX	N-P5DB; AAC64083.		
PT	Novel nucleic acids encoding MiRP1, MiRP2 and MiRP3, useful for		
PT	diagnosing and treating ion channel disorders, especially long QT		
XX	syndrome -		
XX	Claim 56; Page -; 132pp; English.		
CC	The invention relates to novel ion channel proteins related to		
CC	KCNK1 (MinK) and to nucleic acids encoding them. The proteins of		
CC	KCNK1 are human MinK, human MinK2 (MiRP1; AB29593 and AB29598,		
CC	respectively), human MinK3 (KCNE2; AAB29597 and AAB29598,		
CC	respectively). The cDNAs encoding these proteins are given in AAC64071-		
CC	AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier		
CC	potassium channels (I <sub>Kr</sub> ), mutations in which are associated with long		
CC	QT syndrome. The invention also relates to methods of diagnosing long QT		
CC	syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a		
CC	disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic		
CC	nonhuman animals comprising a heterologous ion channel protein gene		
CC	expressing such a gene, and methods of treating long QT syndrome		
CC	using KCNE2 proteins (including mutants), nucleic acids encoding them		
CC	and antibodies against KCNE2 proteins. The methods, antibodies, nucleic		
CC	acids, and proteins may be used for diagnosing or treating ion channel		
CC	disorders, especially long QT syndrome. Transgenic animals comprising		

CC	KCNK2 and HENG are useful for testing anti-long QT syndrome drugs.
CC	The present sequence represents a mutant human KCNE2 (MiRP1)
CC	specifically claimed for use in diagnostic and drug screening methods of
CC	the invention.
CC	The present sequence is not shown in the specification, but is
CC	derived from the wild type human KCNE2 protein sequence shown on page
CC	119.
XX	
SQ	Sequence 123 AA;
Dy	Query Match 99.5%; Score 629; DB 21; Length 123;
Dy	Best Local Similarity 99.2%; Pred. No. 4.4e-67;
Dy	Matches 122; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
OY	1 MSTLSNFFQLDEDYDFRRFTTDMNRONTATAGALAKVDANFNYYVLTLYWIMCF 60
OY	1 MSTLSNFFLEDDYFRRFTTDMNRONTATAGALAKVDANFNYYVLTLYWIMCF 60
Dy	61 SFTIIVALLIVSYVKSRKHSHNDPHOTIVEDEMRKYSQILINBSKAITHEHTGGAQPK 120
Dy	61 SFTIIVALLIVSYVKSRKHSHNDPHOTIVEDEMRKYSQILINBSKAITHEHTGGAQPK 120
OY	121 MSP 123
OY	121 MSP 123
Dy	121 MSP 123
RESULT 12	
ID	AEE22094
AC	AEE22094 standard; Protein; 123 AA.
AC	AEE22094:
DY	25-JUL-2002 (first entry)
DX	
DX	Human MiRP1 mutant protein (A1515).
KM	Human; Min-K related ion channel protein; MiRP1; ion channel disorder;
KM	KCNK2; Long QT syndrome; LQTS; cardiac arrhythmia; mutant; mulein.
OS	Homo sapiens.
XX	
XX	Key Location/Qualifiers
FT	Misc-difference 1 /note= "Wild type Ala substituted with Val"
XX	
XX	MO200222875-A2.
PX	
PD	21-MAR-2002.
PE	11-SEP-2001; 2001MO-US28332.
PR	11-SEP-2000; 2000US-231571P.
XX	(UYVA ) UNITY YALE.
XX	
XI	Goldstein SAN;
XX	
DR	WPI: 2002-362360/39.
DR	N-P50B; RAD35163.
PT	Novel gene encoding Min-K related ion channel protein subunit and
PT	polymorphisms in this gene associated with antibiotic-induced long QT
PT	syndrome -
PS	Claim 1; Page 42; 49pp; English.
XX	
XX	The present invention relates to novel KCNE2 genes encoding Min-K related
XX	ion channels and polymorphisms therein. The invention also relates to
XX	associated with ion channel disorders including antibiotic-induced long
XX	QT syndrome (LQTS). Detecting a mutation at amino acid positions 8, 54,
XX	57 or 116 of MiRP1 polypeptide or a mutation at a nucleotide position
XX	encoding the amino acid positions is useful for diagnosing the presence

CC potassium channels (I-KR), mutations in which are associated with long

XX  
XX  
New] and the and WINN "COEN for





GenCore version 5.1.4\_p5\_4578  
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OK nucleic - nucleic search, using sw model

Run on: May 21, 2003, 19:45:19 ; Search time 596.902 Seconds

(without alignments)  
10011.921 Million cell updates/sec

Title: us-09-550-163-1\_COPY\_74\_442

Sequence: 1 atgtactactatcatcaatt.....ctgggttcaaaatgccc 369

Scoring table: IDENTITY\_NUC  
Gapop 10.0 ; Gapext 1.0

Searched: 16154066 seqs, 809774376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database :

Listing first 45 summaries

ESR:\*  
1: em.estab:\*  
2: em.esthum:\*  
3: em.estlin:\*  
4: em.estlin:\*  
5: em.estlin:\*  
6: em.estlin:\*  
7: em.estlin:\*  
8: em.estlin:\*  
9: gb.est1:\*  
10: gb.est2:\*  
11: gb.est3:\*  
12: gb.est3:\*  
13: gb.est3:\*  
14: gb.est5:\*  
15: em.esthum:\*  
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17: gb.est3:\*  
18: em.gss.hum:\*  
19: em.gss.hum:\*  
20: em.gss.hum:\*  
21: em.gss.hum:\*  
22: em.gss.hum:\*  
23: em.gss.hum:\*  
24: em.gss.hum:\*  
25: em.gss.other:\*  
26: em.gss.pro:\*  
27: em.gss.pro:\*

Prod. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result	No.	Score	Query	Length	DB	ID	Description
c	1	354.8	96.2	403	12	BS208163	RS272654
	2	333	90.2	410	9	A1962650	A1962650
	3	333	90.2	429	9	A1654552	W446012.x
	4	328.8	89.1	391	9	A1339609	q424077.x
	5	325	88.1	372	9	A1246239	q129904.x
	6	266.6	72.2	1691	11	AK008619	AK008619

Result	No.	Score	Query	Length	DB	ID	Description
c	1	354.8	96.2	403	12	BS208163	RS272654
	2	333	90.2	410	9	A1962650	A1962650
	3	333	90.2	429	9	A1654552	W446012.x
	4	328.8	89.1	391	9	A1339609	q424077.x
	5	325	88.1	372	9	A1246239	q129904.x
	6	266.6	72.2	1691	11	AK008619	AK008619

```

SOURCE          1. 603
                 /organism="Homo sapiens"
                 /db_xref="taxon:9606"
                 /clone.lib="Athensys RAGE Library"
                 /cell_line="H1080"
                 /note="See 'Creation of Genome-wide Protein Expression
                 Libraries using Random Activation of Gene Expression',
                 Nature Biotechnology, in press. Note that even though the
                 method was used, these sequences were not necessarily
                 expressed in H1080 under normal circumstances."

BASE COUNT      222 a 172 c 171 g 238 t
ORIGIN

Query Match
Base Count      96 2s; Score 354.8; DB 12; Length 803;
Similarity      96.2s; Pred. No. 3.3e-92;
Matches 367; Conservative 0; Mismatches 2; Indels 1; Gaps 1;

OY 1 AGTCGACTTACTTACCAATTTTCACAGACGCTGGAGAGAGCTCCG-AGAGATTTTTAT 59
DB 688 AGTCGACTTACTTACCAATTTTCACAGACGCTGGAGAGAGCTCCGAGAGATTTTTAT 629
OY 60 TACTTATATGAGCATATGGCGCGGACGAGACAGACAGCTGACGAGAGGCTCTCCAGCCCA 119
DB 628 TACTTATATGAGCATATGGCGCGGACGAGACAGACAGCTGACGAGAGGCTCTCCAGCCCA 569
OY 120 AGTATGCTGAGACCTTCTACTATGTCATGCTGACGCTGGAGATGATGGAATTTT 179
DB 568 AGTATGCTGAGACCTTCTACTATGTCATGCTGACGCTGGAGATGATGGAATTTT 509
OY 180 CTCTTCATCATGCTGACGCTGCTGCTGATGCTGATGCTGATGATGATGATGATGAT 239
DB 508 CTCTTCATCATGCTGACGCTGCTGCTGATGCTGATGCTGATGATGATGATGATGAT 449
OY 240 CAATGACCCCTTACACACGCTGATGCTGATGCTGATGCTGATGCTGATGCTGATGCTG 299
DB 448 CAATGACCCCTTACACACGCTGATGCTGATGCTGATGCTGATGCTGATGCTGATGCTG 389
OY 300 CTTCGATCTGAGAGATGAGAGAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 359
DB 388 CTTCGATCTGAGAGATGAGAGAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 329
OY 360 AATGTCCTCC 369
DB 328 AATGTCCTCC 319

RESULT 2
LOCUS          A1962650          410 bp      mRNA      linear      EST 08-MAR-2000
DEFINITION    w42603.x1 NCI-CGAP GC6 Homo sapiens cDNA clone IMAGE:2473948 3'
               smaller to SW:MIK; HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM
               CHANNEL PROTEIN 1; mRNA sequence.
ACCESSION     A1962650
VERSION       A1962650.1 GI:5755363
KEYWORDS      EST.
SOURCE        human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
NCBI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cga@btl-femail.nih.gov
Tissue Procurement: Christopher A. Woshluk, M.D., Ph.D., Michael
K. Chinnai, M.D., Ph.D.
NCBI-CCAP Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center

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FEATURES
SOURCE
                 1. 410
                 /organism="Homo sapiens"
                 /db_xref="taxon:9606"
                 /clone.lib="NCI-CGAP GC6"
                 /tissue.type="pooled germ cell tumors"
                 /lab_host="DHL03"
                 /note="Vector: pT730-Pac (Pharmacia) with a modified
                 polylinker; Site 1: Not I; Site 2: Eco RI; plasmid DNA
                 from the normalized library NCI-CGAP GC6 was prepared, and
                 six clones were made in vitro. Following the purification
                 reaction, the driver was PCR-amplified cDNA from a pool
                 of 5,000 clones made from the same library (clonides
                 1257096-1258631, 1459064-1470933, and 1475592-1476743).
                 Subtraction by Bento Soares and M. Fatima Bonaldo."
BASE COUNT      120 a 93 c 93 g 104 t
ORIGIN

Query Match
Base Count      90 2s; Score 333; DB 9; Length 410;
Similarity      100.0s; Pred. No. 5.4e-86;
Matches 333; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 37 GAGCTCTTCGAGAGATTTTATCTATGATGACATTTGGCGCGAGACAGACGCT 95
DB 18 GAGCTCTTCGAGAGATTTTATCTATGATGACATTTGGCGCGAGACAGACGCT 77
OY 97 GAGCGAGAGGCGCTCCAGCAAGGATGATGATGATGATGATGATGATGATGATGATG 156
DB 78 GAGCGAGAGGCGCTCCAGCAAGGATGATGATGATGATGATGATGATGATGATGATG 137
OY 157 CTCATGCTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 216
DB 138 CTCATGCTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 197
OY 217 AATTCGAGAGAGGAGAGATGATGATGATGATGATGATGATGATGATGATGATG 276
DB 198 AATTCGAGAGAGGAGAGATGATGATGATGATGATGATGATGATGATGATGATG 257
OY 277 CAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 336
DB 258 CAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 317
OY 337 AACCATGCTGCGGCTGCGGCTGCGGCTGCGGCTGCGGCTGCGGCTGCGGCTGCGG 369
DB 318 AACCATGCTGCGGCTGCGGCTGCGGCTGCGGCTGCGGCTGCGGCTGCGGCTGCGG 350

RESULT 3
LOCUS          A1654552          429 bp      mRNA      linear      EST 17-DEC-1999
DEFINITION    w448d12.x1 NCI-CGAP GC6 Homo sapiens cDNA clone IMAGE:2308895 3'
               smaller to SW:MIK; HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM
               CHANNEL PROTEIN 1; mRNA sequence.
ACCESSION     A1654552
VERSION       A1654552.1 GI:4738531
KEYWORDS      EST.
SOURCE        human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
NCBI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.

```

Email: [cgapbs-remail.nih.gov](mailto:cgapbs-remail.nih.gov)  
 Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.  
 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonalido, Ph.D.  
 Library Arrayed by: Greg Lennon, Ph.D.  
 DNA Sequencing: J. Michael Winkler, University of Michigan Sequencing Center  
 Found through the I.M.A.G.E. Consortium/LNL at: [www-bio.lnlnl.gov/obrp/image/image.html](http://www-bio.lnlnl.gov/obrp/image/image.html)  
 Insert Length: 771 Std Error: 0.00  
 Seq primer: -40bp from Glbco  
 High quality sequence stop: 411.  
 Location/Qualifiers  
 1..429  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone="IMAGE:2308895"  
 /clone\_1lb="NCI-CGAP-GC6"  
 /tissue\_type="pooled germ cell tumors"  
 /note="Vector: p773D-pac (Pharmacia) with a modified polylinker. Site 1: Not I; Site 2: Eco RI; Plasmid DNA from the normalized library NCI-CGAP-GC4 was prepared, and 88 circles were made in vitro. Following BAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool 1237090-125851 (489441/489442/489443)."  
 Subtraction by Bento Soares and M. Fatima Bonalido.

BASE COUNT 127 a 100 c 97 g 104 t 1 others

Query Match Best Local Similarity 90.24; Score 333; DB 9; Length 429;  
 Best Local Similarity 100.08; Pred. No. 5.5e-86;  
 Matches 333; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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OY 37 GAGCTCTCGAAGAGATTTTATTACTATATGACCAATTGGCGCCGACACACACAGCT 96
    |||||||
DB 12 GAGCTCTCGAAGAGATTTTATTACTATATGACCAATTGGCGCCGACACACACAGCT 71
OY 97 GAGCAGAGGCGCTCCAGACCCAAAGTGTGAGAGAACTTCTACTATGATTCATCTGTAC 156
    |||||||
DB 72 GAGCAGAGGCGCTCCAGACCCAAAGTGTGAGAGAACTTCTACTATGATTCATCTGTAC 131
OY 157 CTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 216
    |||||||
DB 132 CTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 191
OY 217 AATTCGAAAGAGGCGACACCTCATGACCCCTCCACGACATCTGTAGAGACATGG 276
    |||||||
DB 192 AATTCGAAAGAGGCGACACCTCATGACCCCTCCACGACATCTGTAGAGACATGG 251
OY 277 CAGGAAAGTACAGAGCAATCTGATATGAGAAAGTACAGAGCCACATCCATGAG 336
    |||||||
DB 252 CAGGAAAGTACAGAGCAATCTGATATGAGAAAGTACAGAGCCACATCCATGAG 311
OY 337 AACTGTGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCG 369
    |||||||
DB 312 AACTGTGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCG 344

```

RESULT 4  
 A1336609 301 bp mRNA linear EST 29-DEC-1998  
 DEFINITION 37 similar to SW:MIK\_HUMAN\_P13382 ISK SLOW VOLTAGE GATED POTASSIUM CHANNEL PROTEIN ; mRNA sequence.  
 ACCESSION A1336609  
 VERSION A1336609.1 GI:4076536  
 KEYWORDS EST.  
 SOURCE Human.  
 ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 391)  
 AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.  
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index  
 JOURNAL NCI-CGAP (1997)  
 COMMENT (Cont.) Robert Strumberg, Ph.D.  
 Email: [cgapbs-remail.nih.gov](mailto:cgapbs-remail.nih.gov)  
 This clone is available royalty-free through LNL; contact the IMAGE Consortium ([info@image.lnlnl.gov](mailto:info@image.lnlnl.gov)) for further information.  
 Seq primer: -40bp from Glbco  
 High quality sequence stop: 380.  
 Location/Qualifiers  
 1..391  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone="IMAGE:1931516"  
 /clone\_1lb="Soares NIHMPu.S1"  
 /tissue\_type="Pooled human melanocyte, fetal heart, and pregnant uterus"  
 /note="Organ: mixed (see below); Vector: p773D-pac (Pharmacia) with a modified polylinker. Site 1: Not I; Site 2: Eco RI; Equal amounts of plasmid DNA from three normalized libraries (melanocyte 2bhm, pregnant uterus Nbhpv, and fetal heart Nbhp19v) were mixed, and 88 circles were made in vitro. Following BAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same 3 libraries. The pools consisted of I.M.A.G.E. clones 260232-265223, 340488-345479, and 484488-489479."

BASE COUNT 119 a 93 c 93 g 86 t

Query Match Best Local Similarity 89.14; Score 328.8; DB 9; Length 391;  
 Best Local Similarity 99.48; Pred. No. 8.8e-85;  
 Matches 330; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

```

OY 38 ACGCTCTCGAAGAGATTTTATTACTATATGACCAATTGGCGCCGACACACAGCTG 97
    |||||||
DB 3 ACGCTCTCGAAGAGATTTTATTACTATATGACCAATTGGCGCCGACACACAGCTG 62
OY 96 AGGAGAGGCGCTCCAGACCCAAAGTGTGAGAGAACTTCTACTATGATTCATCTGTAC 157
    |||||||
DB 63 AGGAGAGGCGCTCCAGACCCAAAGTGTGAGAGAACTTCTACTATGATTCATCTGTAC 122
OY 156 TCAATGTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 217
    |||||||
DB 123 TCAATGTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 182
OY 218 AATTCGAAAGAGGCGACACCTCATGACCCCTCCACGACATCTGTAGAGACATGG 277
    |||||||
DB 183 AATTCGAAAGAGGCGACACCTCATGACCCCTCCACGACATCTGTAGAGACATGG 242
OY 278 AGGAAAGTACAGAGCAATCTGATATGAGAAAGTACAGAGCCACATCCATGAG 337
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DB 243 AGGAAAGTACAGAGCAATCTGATATGAGAAAGTACAGAGCCACATCCATGAG 302
OY 338 AACTGTGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCG 369
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DB 303 AACTGTGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCGGTGCG 334

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RESULT 5  
 A1246239 312 bp mRNA linear EST 28-JAN-1999  
 DEFINITION 37 similar to SW:MIK\_HUMAN\_P13382 ISK SLOW VOLTAGE GATED POTASSIUM CHANNEL PROTEIN ; mRNA sequence.  
 ACCESSION A1246239  
 VERSION A1246239.1 GI:3841636







Oy	241	AATGACCCCTCACCAACCAGTAAATTGTGAGAGACTGGCAGGGAAAAGCTPACAAGAGCAAAATTC	300
Oy	277	TGGAGACCGCGGCACACAGATGNCCTGATGCTGGAGGATTTGGCAGCAAAGATACGAGCATC	336
Oy	301	TGGAATCTCAGAACAGATCCAGATCCAGATCCGAGCAACCACTTGTCGGCTGGGTTCANA	360
Dy	337	TGAGATCTCGAAGATCTCCAGGCCATCATCATCATATTAACCTTGAGGGGAGGGGTTCACA	396
Oy	361	ATGTCTCCCC	369
Dy	397	MTGTACCC	405

RESULT 8  
Bg261965

LOCUS Bg261965 1003 bp mRNA linear EST 13-FEB-2003  
DEFINITION 6127377461.NIH\_MGC\_94 Mus musculus cDNA clone IMAGE:4481325 5,'  
VERSION 1.0 sequence.  
ACCESSION Bg261965  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus.  
REMARKS Mammalia; Eumetazoa; Chordata; Crustacea; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
REFERENCE 1 (bases 1 to 1003)  
TITLE NIH-MGC http://mgc.nci.nih.gov/.  
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL Unpublished (1999)  
COMMENT Contact: Robert Strausberg, Ph.D.  
Contact: David Lipman, Ph.D.  
Tissue Procurement: The CCRB Laboratory  
CDNA Library Preparation: Life Technologies, Inc.  
DNA Sequencing By: Incyte Genomics, Inc.  
Clone Distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/ILMM at:  
http://www.imm.kit.edu/  
Plate: ILMM0316 ROW: g column: 22  
High quality sequence stop: 535.

Location/Qualifiers  
1..1003

FEATURES  
source

	Query Match	65.6% Best Local Similarity	Score: 242; ID 52; 81.7% Prec. No. 2; Le 19;	length 1003;
	Matches 304; Conservative	0; Mismatches 65;	Indels 3;	Gaps 2;
07	1 ATGTCACCTTATTCACAAATTCACACAGAGGTGTGAAACGCTCTCCAGAGGATTTTTTATTTT	60		
Db	91 ATGTCACCTTATTCACAAATTCACACAGAGGTGTGAAACGCTCTCCAGAGGATTTTTTATTTT	150		
07	61 ACTTATGAGGACATATTCGCAATTTTGACACAGACAGCTGATGAGATGCGCTTCATAGCCGAA	120		
Db	151 ACTTATGAGGACATATTCGCAATTTTGACACAGACAGCTGATGAGATGCGCTTCATAGCCGAA	210		
07	121 GTTATGCTGTGAGAACTCTTCATCATGTGACATCCCTGTACCTCATAGGTGATGTGAAAGTTC	180		
Db	211 GTTATGCTGTGAGAACTCTTCATCATGTGACATCCCTGTACCTCATAGGTGATGTGAAAGTTC	370		

Qy	181	TCCTCATCATCTGGGCGCATCTGGGTATACACTGTGAAATTCAGAGAGAGAGAGACTTC	240
Db	271	TGCTTCATCTGGTGGCGCATCTCGGTATACACAGGGAAATTCAGAGAGAGAGACTTC	330
Qy	241	CAACACCCCTACACACACATATGTAAGAGACTGGCAGGAAAGTACAGAGCCAAATC	300
Db	331	CAGCAGCCGCTGACACACACATATGTAAGAGACTGGCAGGAAAGTACAGAGCCAAATC	390
Qy	301	TTGATATCTATAGAAAGACGAA--GCGCCACATCCATGAGAGAAACT--TGAGGTCTGGGATTC	357
Db	391	CTGATCTCGGAAGACCTCCAAAGGCGACCCATCCATGAGAGAACTTGAGGAGGCGAGCGGGTTC	450
Qy	358	AAATATGTCCGCC	369
Db	451	ACGATGTACCC	462

RESULT 9			
LOCUS	Bg221966		
DEFINITION	Bg221966	746 bp	mRNA
VERSION	BS221966.1		Linear
KEYWORDS	EST.		EST 21-APR-2001
SOURCE	human.		
ORGANISM	Homo sapiens		
REFERENCE	Bukaryaga, Meszaca, Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eularchia; Primates; Cetartihud; Homalodonta; Homo. 1 (bases 1 to 746)		
AUTHORS	Dair, J., Gordon, A., Sherr, P., Rundlett, S., Jackson, P. D., Berry, R., Gaitanaris, G., Chittala, J., Smith, J. A., Hwang, J., Lerner, D., Costanzo, D., McMilligott, K., Boyer, S., Myers, R., Smith, J., Veloso, N., Kika, A., Hess, J., Colburn, K., Lo, K., Offenbacher, J., Danzig, J., and Dincer, M.		
TITLE	Creation of genome-wide protein expression libraries using random activation of gene expression		
JOURNAL	Nat. Biotechnol.	19 (5),	440-445 (2001)
PMID	11227151		
COMMENT	Scott J. Cain		
ALTERNATIVE	Athersys, Inc.		
COMMENT	3201 Carnegie Ave., Cleveland, OH 44115, USA		
	Tel.: 216 431 9900		
	Fax: 216 361 9596		
	Email: sca@atersys.com		
	High quality sequence atop: 547.		

Query Match	Local Similarity	Score	DB	Length
1	97.2%	177.8	12	746
2	97.2%	177.8	12	746
3	97.2%	177.8	12	746
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25	97.2%	177.8	12	746
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77	97.			







Db 1 CTTGAATCTAGAGATCGAGGCCACCATCCATGAGAACATTGGTCCGGCTGGGTCAA 60  
Oy 360 AATGTCCCC 369  
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Db 61 AATGTCCCC 70

Search completed: May 21, 2003, 21:37:55  
Job time : 600.902 secs

GenCore version 5.1.4-p5.4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using bw model

Run on: May 21, 2003, 20:27:39 (Search time 50.6076 Seconds  
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9638.006 Million cell updates/sec

Title: US-09-550-163-1-COPY\_74\_442

Perfect score: 369  
Sequence: 1 agcctacttaccatc.....ctgggtcctaaatgcctccc 369

Scoring table: IDENTITY NTC  
Gapco 10.0 , Gapext 1.0

Searched: 828747 sets, 660231138 residues

Total number of hits satisfying chosen parameters: 1657494

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database : Listing first 45 summaries

Database :	Published Applications NA *
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the total score distribution, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	369	100.0	372	10	US-09-864-761-33139 Sequence 33139, A
2	369	100.0	732	9	US-10-000-1518-5 Sequence 5, Appl1
3	369	100.0	113604	9	US-10-227-195A-1 Sequence 1, Appl1
4	369	100.0	113604	9	US-10-227-195A-1 Sequence 2, Appl1
5	369	100.0	113604	9	US-10-227-195A-1 Sequence 3, Appl1
6	369	100.0	113604	9	US-10-227-195A-1 Sequence 4, Appl1
7	369	100.0	113604	9	US-10-227-195A-1 Sequence 5, Appl1
8	369	100.0	113604	9	US-10-227-195A-1 Sequence 6, Appl1
9	369	100.0	113604	9	US-10-227-195A-1 Sequence 7, Appl1
10	369	100.0	113604	9	US-10-227-195A-1 Sequence 8, Appl1
11	369	100.0	113604	9	US-10-227-195A-1 Sequence 9, Appl1
12	369	100.0	113604	9	US-10-227-195A-1 Sequence 10, Appl1
13	369	100.0	113604	9	US-10-227-195A-1 Sequence 11, Appl1
14	369	100.0	113604	9	US-10-227-195A-1 Sequence 12, Appl1
15	369	100.0	113604	9	US-10-227-195A-1 Sequence 13, Appl1
16	369	100.0	113604	9	US-10-227-195A-1 Sequence 14, Appl1
17	369	100.0	113604	9	US-10-227-195A-1 Sequence 15, Appl1
18	369	100.0	113604	9	US-10-227-195A-1 Sequence 16, Appl1
19	369	100.0	113604	9	US-10-227-195A-1 Sequence 17, Appl1
20	369	100.0	113604	9	US-10-227-195A-1 Sequence 18, Appl1

## ALIGNMENTS

20	36	9.8	2579	10	US-09-822-830A-30	Sequence 30, Appl1
21	33.4	9.1	4104	9	US-09-992-598-277	Sequence 277, App
22	33.4	9.1	4104	9	US-09-989-293A-277	Sequence 277, App
23	33.4	9.1	4104	9	US-09-989-735-277	Sequence 277, App
24	33.4	9.1	4104	9	US-09-990-444-277	Sequence 277, App
25	33.4	9.1	4104	9	US-09-989-730-277	Sequence 277, App
26	33.4	9.1	4104	9	US-09-991-183-277	Sequence 277, App
27	33.4	9.1	4104	9	US-09-993-687-277	Sequence 277, App
28	33.4	9.1	4104	9	US-09-989-734-277	Sequence 277, App
29	33.4	9.1	4104	9	US-10-028-072-449	Sequence 449, App
30	33.4	9.1	4104	9	US-09-997-653-277	Sequence 277, App
31	33.4	9.1	4104	9	US-09-997-657-277	Sequence 277, App
32	33.4	9.1	4104	9	US-10-121-069-449	Sequence 449, App
33	33.4	9.1	4104	9	US-10-140-470-449	Sequence 449, App
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42	33.4	9.1	4104	9	US-10-175-746-449	Sequence 449, App
43	33.4	9.1	4104	9	US-10-227-884-157	Sequence 157, App
44	33.4	9.1	4104	9	US-09-990-711-277	Sequence 277, App
45	33.4	9.1	4104	9	US-10-137-865-449	Sequence 449, App

## RESULTS

1	US-09-864-761-33139	Sequence 33139, Application US/09864761
2	Patent No. US20020048763A1	
3	GENERAL INFORMATION:	
4	APPLICANT: Penn, Sharon G.	
5	APPLICANT: Hema, David A.	
6	APPLICANT: Hema, David A.	
7	APPLICANT: Chem, Wensheng	
8	TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL F	
9	FILE REFERENCE: Aeonica-X-1	
10	CURRENT APPLICATION NUMBER: US/09/864,761	
11	PRIOR APPLICATION NUMBER: US 60/180,312	
12	PRIOR FILING DATE: 2000-02-04	
13	PRIOR APPLICATION NUMBER: US 60/207,456	
14	PRIOR FILING DATE: 2000-05-26	
15	PRIOR APPLICATION NUMBER: US 09/632,366	
16	PRIOR FILING DATE: 2000-08-03	
17	PRIOR APPLICATION NUMBER: US 60/246,316	
18	PRIOR FILING DATE: 2000-09-27	
19	PRIOR APPLICATION NUMBER: PCT/US01/00666	
20	PRIOR FILING DATE: 2001-01-30	
21	PRIOR APPLICATION NUMBER: PCT/US01/00666	
22	PRIOR FILING DATE: 2001-01-30	
23	PRIOR APPLICATION NUMBER: PCT/US01/00666	
24	PRIOR FILING DATE: 2001-01-30	
25	PRIOR APPLICATION NUMBER: PCT/US01/00666	
26	PRIOR FILING DATE: 2001-01-30	
27	PRIOR APPLICATION NUMBER: PCT/US01/00666	
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44	PRIOR FILING DATE: 2001-01-30	
45	PRIOR APPLICATION NUMBER: PCT/US01/00666	

PRIOR APPLICATION NUMBER: PCT/US01/00670  
 PRIOR FILING DATE: 2000-06-30, EST. HUMAN HIT: 0.60/234,687  
 PRIOR APPLICATION NUMBER: US 09/608,408  
 PRIOR FILING DATE: 2000-06-30  
 PRIOR APPLICATION NUMBER: US 09/774,203  
 PRIOR FILING DATE: 2001-01-29  
 NUMBER OF SEQ ID NOS: 43117  
 SEQUENCE LENGTH: 1115  
 SEQ ID NO: 3139  
 LENGTH: 372  
 TYPE: DNA  
 ORGANISM: Homo sapiens  
 FEATURE: MAP TO AF000120.1  
 OTHER INFORMATION: EXPRESSED IN HL100 SIGNAL: 0.67  
 OTHER INFORMATION: EST. HUMAN HIT: A155452.1, EVALUATE 0.00e+00  
 OTHER INFORMATION: SWISSPROT HIT: Q9Y6J6, EVALUATE 8.00e-67  
 OTHER INFORMATION: NCBI HIT: g11526220, EVALUATE 0.00e+00  
 US-09-564-761-3139

Query Match 100.0%; Score 369; DB 10; Length 732;  
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 Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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 OY 181 TCTTTCATCATGCTGGCAATCTGTGTGACAGCTGTAAATCCAGAAAGGAAACATGTC 240  
 DB 181 TCTTTCATCATGCTGGCAATCTGTGTGACAGCTGTAAATCCAGAAAGGAAACATGTC 240  
 OY 241 AATGACCCCTACACAGTACATTTGTAGAGAGATGGCGAGGAAATGTCAAGACCCAAATC 300  
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 OY 301 TTGAATCTAGAGATTCAGAGGCCACATCCATGATAGAACATTTGGTGGCGTTTCAA 360  
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 OY 361 ATGTCCGCC 369  
 DB 361 ATGTCCGCC 369

RESULT 2  
 US-10-000-151B-5  
 Sequence 5, Application US/10000151B  
 Publication No. US20030013136A1  
 GENE: INFORMATION:  
 ORGANISM: Homo sapiens  
 APPLICANT: Joffrey R.  
 TITLE OF INVENTION: HUMAN KCI REGULATION OF HERG POTASSIUM CHANNEL BLOCK  
 FILE REFERENCE: Vanderbilt Ref No. US20030013136A1 V01020; Attorney Docket No. US2003  
 CURRENT FILING DATE: 05/10/00, 151B  
 NUMBER OF SEQ ID NOS: 5  
 SOFTWARE: PatentIn version 3.1  
 SEQ ID NO: 1  
 LENGTH: 732  
 TYPE: DNA  
 ORGANISM: Homo sapiens

US-10-000-151B-5  
 Query Match 100.0%; Score 369; DB 9; Length 732;  
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 Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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 DB 134 ACTTATATGACAAATTTGGCGCAAGAACACAGCTGAGACAGAGCCCTCCAGACCAA 193  
 OY 121 GTTATGCTGAGAACTTCTACTATGTCATCTGTCATGCTCATGCTGATGATTTGAAATGTC 180  
 DB 194 GTTATGCTGAGAACTTCTACTATGTCATCTGTCATGCTCATGCTGATGATTTGAAATGTC 253  
 OY 181 TCTTTCATCATGCTGGCAATCTGTGTGACAGCTGTAAATCCAGAAAGGAAACATGTC 240  
 DB 254 TCTTTCATCATGCTGGCAATCTGTGTGACAGCTGTAAATCCAGAAAGGAAACATGTC 313  
 OY 241 AATGACCCCTACACAGTACATTTGTAGAGAGATGGCGAGGAAATGTCAAGACCCAAATC 300  
 DB 314 AATGACCCCTACACAGTACATTTGTAGAGAGATGGCGAGGAAATGTCAAGACCCAAATC 373  
 OY 301 TTGAATCTAGAGATTCAGAGGCCACATCCATGATAGAACATTTGGTGGCGTTTCAA 360  
 DB 374 TTGAATCTAGAGATTCAGAGGCCACATCCATGATAGAACATTTGGTGGCGTTTCAA 433  
 OY 361 ATGTCCGCC 369  
 DB 434 ATGTCCGCC 442

RESULT 3  
 US-10-227-195A-1  
 Sequence 1, Application US/10227195A  
 Publication No. US2003007653A1  
 GENE: INFORMATION:  
 ORGANISM: Homo sapiens  
 APPLICANT: Cox, David  
 TITLE OF INVENTION: Haplotype structure of chromosome 21  
 FILE REFERENCE: 101001  
 CURRENT APPLICATION NUMBER: US/10/227,195A  
 CURRENT FILING DATE: 2002-11-18  
 NUMBER OF SEQ ID NOS: 2  
 SOFTWARE: TestSeq for Windows version 4.0  
 SEQ ID NO: 1  
 LENGTH: 113604  
 TYPE: DNA  
 ORGANISM: Human  
 FEATURE: misc-feature  
 NAME/REF: US/10/227,195A, 36972, 76921, 81512, 88727  
 OTHER INFORMATION: n = 6 or C  
 US-10-227-195A-1

Query Match 100.0%; Score 369; DB 9; Length 113604;  
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 Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATGCTCTACTTATTCACATTCACACAGACGCTGAGACGCTTCGCAAGAGATTTTATT 60  
 DB 17476 ATGCTCTACTTATTCACATTCACACAGACGCTGAGACGCTTCGCAAGAGATTTTATT 17535  
 OY 61 ACTTATATGACAAATTTGGCGCAAGAACACAGCTGAGACAGAGCCCTCCAGACCAA 120  
 DB 17536 ACTTATATGACAAATTTGGCGCAAGAACACAGCTGAGACAGAGCCCTCCAGACCAA 17595  
 OY 121 GTTATGCTGAGAACTTCTACTATGTCATCTGTCATGCTCATGCTGATGATTTGAAATGTC 180  
 DB 17596 GTTATGCTGAGAACTTCTACTATGTCATCTGTCATGCTCATGCTGATGATTTGAAATGTC 17655



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OY 181 TCTTTCATCATGTCGTGGCCACTGCTGTGAGCACTGTGAATTCAGAGAGCGGAGAACATCTCC 240
|||||
Db 17656 TCTTTCATCATGTCGTGGCCACTGCTGTGAGCACTGTGAATTCAGAGAGCGGAGAACATCTCC 17715
OY 241 AATGACCCCTTACACACAGTACATGTTGTGAGAGCGGCGAGAAAATGACAGACCCAAATC 300
|||||
Db 17716 AATGACCCCTTACACACAGTACATGTTGTGAGAGCGGCGAGAAAATGACAGACCCAAATC 17775
OY 301 TTGAATCTTGAAGAAATGAGAGGCGACACATCCATGAAACATTTGTGCTGGGTGCTTAA 360
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Db 17776 TTGAATCTTGAAGAAATGAGAGGCGACACATCCATGAAACATTTGTGCTGGGTGCTTAA 17835
OY 361 ATGCTCCGCC 369
|||||
Db 17836 ATGCTCCGCC 17844

RESULT 4
US-10-227-195A-2
Query Match 100.0%; Score 369; DB 9; Length 113604;
Best Local Similarity 100.0%; Pred. No. 3,6e-114;
Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

GENERAL INFORMATION:
PUBLICATION: Appl US2003007633A1
APPLICANT: Cox, David
APPLICANT: Arnold, Deana
TITLE OF INVENTION: Haplotype structure of chromosome 21
FILE REFERENCE: 103001
CURRENT FILING DATE: 2001-01-30
NUMBER OF SEQ ID NOS: 2
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO: 2
LENGTH: 113604
TYPE: DNA Human
US-10-227-195A-2

Query Match 100.0%; Score 369; DB 9; Length 113604;
Best Local Similarity 100.0%; Pred. No. 3,6e-114;
Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATGCTACTTATTCATATTCACACACAGCTGTGAGAGAGCTTTCTCGAAGATTTTAT 60
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Db 17476 ATGCTACTTATTCATATTCACACACAGCTGTGAGAGAGCTTTCTCGAAGATTTTAT 17535
OY 61 ACTTAATATGATGAATGTGGCGCAGAAACAACAGCTGTGAGAGAGCGCTCCACACCAA 120
|||||
Db 17536 ACTTAATATGATGAATGTGGCGCAGAAACAACAGCTGTGAGAGAGCGCTCCACACCAA 17595
OY 121 GTTGATGTGAAGAACTTACTATGTATCTGTAATCTGTAATCTGTAATCTGTAATCTGTA 180
|||||
Db 17596 GTTGATGTGAAGAACTTACTATGTATCTGTAATCTGTAATCTGTAATCTGTAATCTGTA 17655
OY 181 TCTTTCATCATGTCGTGGCCACTGCTGTGAGCACTGTGAATTCAGAGAGCGGAGAACATCTCC 240
|||||
Db 17656 TCTTTCATCATGTCGTGGCCACTGCTGTGAGCACTGTGAATTCAGAGAGCGGAGAACATCTCC 17715
OY 241 AATGACCCCTTACACACAGTACATGTTGTGAGAGCGGCGAGAAAATGACAGACCCAAATC 300
|||||
Db 17716 AATGACCCCTTACACACAGTACATGTTGTGAGAGCGGCGAGAAAATGACAGACCCAAATC 17775
OY 301 TTGAATCTTGAAGAAATGAGAGGCGACACATCCATGAAACATTTGTGCTGGGTGCTTAA 360
|||||
Db 17776 TTGAATCTTGAAGAAATGAGAGGCGACACATCCATGAAACATTTGTGCTGGGTGCTTAA 17835
OY 361 ATGCTCCGCC 369
|||||
Db 17836 ATGCTCCGCC 17844

RESULT 5
US-09-864-761-20233
Query Match 84.6%; Score 312; DB 10; Length 312;
Best Local Similarity 100.0%; Pred. No. 3,4e-96;
Matches 312; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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? Patent No. US20020048763A1
? GENERAL INFORMATION:
? APPLICANT: Penn, Sharon G.
? APPLICANT: Rank, David R.
? APPLICANT: Hanzel, David K.
? APPLICANT: Hanzel, David K.
? TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL F
? FILE REFERENCE: Aeonica-X-1
? CURRENT FILING DATE: US/09/864,761
? PRIOR APPLICATION NUMBER: US 60/180,312
? PRIOR FILING DATE: 2000-02-04
? PRIOR APPLICATION NUMBER: US 60/207,456
? PRIOR FILING DATE: 2000-05-09
? PRIOR APPLICATION NUMBER: US 09/652,366
? PRIOR FILING DATE: 2000-08-03
? PRIOR APPLICATION NUMBER: GB 24263,6
? PRIOR FILING DATE: 2000-10-04
? PRIOR APPLICATION NUMBER: US 60/236,359
? PRIOR FILING DATE: 2000-09-27
? PRIOR APPLICATION NUMBER: US 60/201,00662
? PRIOR FILING DATE: 2001-01-30
? PRIOR APPLICATION NUMBER: PCT/US01/00667
? PRIOR FILING DATE: 2001-01-30
? PRIOR APPLICATION NUMBER: PCT/US01/00664
? PRIOR FILING DATE: 2001-01-30
? PRIOR APPLICATION NUMBER: PCT/US01/00669
? PRIOR FILING DATE: 2001-01-30
? PRIOR APPLICATION NUMBER: PCT/US01/00665
? PRIOR FILING DATE: 2001-01-30
? PRIOR APPLICATION NUMBER: PCT/US01/00668
? PRIOR FILING DATE: 2001-01-30
? PRIOR APPLICATION NUMBER: PCT/US01/00663
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? PRIOR FILING DATE: 2001-01-30
? PRIOR APPLICATION NUMBER: US 60/234,687
? PRIOR FILING DATE: 2000-09-16
? PRIOR APPLICATION NUMBER: 09/608,408
? PRIOR FILING DATE: 2000-06-30
? PRIOR APPLICATION NUMBER: US 09/774,203
? PRIOR FILING DATE: 2001-01-29
? NUMBER OF SEQ ID NOS: 49117
? SOFTWARE: Anomax Sequence Listing Engine vers. 1.1
? SEQ ID NO: 20233
? LENGTH: 312
? TYPE: DNA
? ORGANISM: Homo sapiens
? FEATURE:
? OTHER INFORMATION: MAP TO AP000052.1
? OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.1
? OTHER INFORMATION: EXPRESSED IN HB1100, SIGNAL = 1.1
? OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.3
? OTHER INFORMATION: EXPRESSED IN BRYAN, SIGNAL = 0.92
? OTHER INFORMATION: EXPRESSED IN BRYAN, SIGNAL = 0.92
? OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 4
? OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.1
? OTHER INFORMATION: EXPRESSED IN HEPA, SIGNAL = 1.2
? OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 0.94
? OTHER INFORMATION: EXPRESSED IN BT74, SIGNAL = 0.88
? OTHER INFORMATION: ESTROKAN HIT: 61246239.1, EVALUO 0.00e+00
? OTHER INFORMATION: ESTROKAN HIT: 61246239.1, EVALUO 0.00e-55
? OTHER INFORMATION: NF HIT: AF302095.1, EVALUO 0.00e+00
US-09-864-761-20233
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Oy 54 TTTTATATCTTATATGCAATTTGCGCCACAGACACACAGCTGACGAGAGAGCCCTTCA 113  
 Db 1 TTTTATATCTTATATGCAATTTGCGCCACAGACACACAGCTGACGAGAGAGCCCTTCA 60  
 Oy 114 ACCCAAAAGTTGATGCTGAGACCTCTCACTATGTCATCTGACCTGACCTGATGATG 173  
 Db 61 ACCCAAAAGTTGATGCTGAGACCTCTCACTATGTCATCTGACCTGACCTGATGATG 120  
 Oy 174 AATGTCCTCTTATCATGATCGTGGCCACCTCTGTCACACCTGTGAATCCAGAGAGCCGA 233  
 Db 121 AATGTCCTCTTATCATGATCGTGGCCACCTCTGTCACACCTGTGAATCCAGAGAGCCGA 180  
 Oy 234 ACATCCCAATGACCCCTACACACAGTATGTCATGAGACCTGACGAGAAAAGTACAGAG 293  
 Db 181 ACATCCCAATGACCCCTACACACAGTATGTCATGAGACCTGACGAGAAAAGTACAGAG 240  
 Oy 294 CCAATCTTGATATGATGAAGATTCGAGAGGCGACCTCCGAGAACTTGGCGCGCTG 353  
 Db 241 CCAATCTTGATATGATGAAGATTCGAGAGGCGACCTCCGAGAACTTGGCGCGCTG 300  
 Oy 354 GTTCAAAAAATGTC 365  
 Db 301 GTTCAAAAAATGTC 312  
 RESULT 6  
 US-09-864-761-3463  
 Sequence 3463, Application US/09864761  
 Patent No. US20020048763A1  
 GENERAL INFORMATION:  
 APPLICANT: Penn, Sharron G.  
 APPLICANT: Smith, David R.  
 APPLICANT: Hazenel, David K.  
 APPLICANT: Chen, Wensheng  
 TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR  
 FILE REFERENCE: A60163-X-1  
 CURRENT FILING DATE: 2001-05-23  
 PRIOR FILING DATE: 2000-02-04  
 PRIOR APPLICATION NUMBER: US 60/180,312  
 PRIOR FILING DATE: 2000-05-26  
 PRIOR APPLICATION NUMBER: US 60/632,366  
 PRIOR FILING DATE: 2000-10-04  
 PRIOR APPLICATION NUMBER: US 60/236,359  
 PRIOR FILING DATE: 2000-09-27  
 PRIOR APPLICATION NUMBER: PCT/US01/00666  
 PRIOR FILING DATE: 2001-01-30  
 PRIOR APPLICATION NUMBER: PCT/US01/00667  
 PRIOR FILING DATE: 2001-05-23  
 PRIOR APPLICATION NUMBER: PCT/US01/00664  
 PRIOR FILING DATE: 2001-01-30  
 PRIOR APPLICATION NUMBER: PCT/US01/00665  
 PRIOR FILING DATE: 2001-01-30  
 PRIOR APPLICATION NUMBER: PCT/US01/00668  
 PRIOR FILING DATE: 2001-01-30  
 PRIOR APPLICATION NUMBER: PCT/US01/00662  
 PRIOR FILING DATE: 2001-01-30  
 PRIOR APPLICATION NUMBER: PCT/US01/00661  
 PRIOR FILING DATE: 2001-01-30  
 PRIOR APPLICATION NUMBER: PCT/US01/00670  
 PRIOR FILING DATE: 2000-09-27  
 PRIOR APPLICATION NUMBER: US 60/234,687  
 PRIOR FILING DATE: 2000-09-21  
 PRIOR APPLICATION NUMBER: US 09/608,408

PRIOR FILING DATE: 2000-06-30  
 PRIOR APPLICATION NUMBER: US 09/774,203  
 PRIOR FILING DATE: 2000-01-29  
 NUMBER OF SEQ IDS: 41175  
 SOFTWARE: Anomax Sequence Listing Engine vers. 1.1  
 SEQ ID NO 3463  
 LENGTH: 450  
 TYPE: DNA  
 ORGANISM: Homo sapiens  
 OTHER INFORMATION: MAP TO AP000052.1  
 OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.1  
 OTHER INFORMATION: EXPRESSED IN BILILOD, SIGNAL = 1.1  
 OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.3  
 OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 0.92  
 OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1  
 OTHER INFORMATION: EXPRESSED IN SPERMATID, SIGNAL = 1.1  
 OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 0.94  
 OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 0.94  
 OTHER INFORMATION: EXPRESSED IN BPA74, SIGNAL = 0.88  
 US-09-864-761-3463  
 Query Match 79.98; Score 295; DB 10; Length 450;  
 Best Local similarity 100.0%; Pctid No. 2.0e-90;  
 Matches 295; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Oy 1 ATGCTACTATTATTCATTTTCACAGACGCTGGAGAGACGCTCTCCAGAGATTATTAT 60  
 Db 156 ATGCTACTATTATTCATTTTCACAGACGCTGGAGAGACGCTCTCCAGAGATTATTAT 215  
 Oy 61 ACTATATGCAATTTGCGCCACAGACACACAGCTGACGAGAGAGCCCTTCCACCCAA 120  
 Db 216 ACTATATGCAATTTGCGCCACAGACACACAGCTGACGAGAGAGCCCTTCCACCCAA 175  
 Oy 121 GTTATGCTGAGACCTTCACTATGTCATCTGACCTGATGATGATGATGATGATG 180  
 Db 276 GTTATGCTGAGACCTTCACTATGTCATCTGACCTGATGATGATGATGATGATG 335  
 Oy 181 TCTTTCATCATGTCGCGCCATCTGTCGACCTGTGAATCCAGAGAGAGGGAACCTCC 240  
 Db 336 TCTTTCATCATGTCGCGCCATCTGTCGACCTGTGAATCCAGAGAGAGGGAACCTCC 395  
 Oy 241 AATGACCCCTTACACACAGTATGTCATGAGACCTGCGAGAAAAGTACAGAGAGCC 295  
 Db 396 AATGACCCCTTACACACAGTATGTCATGAGACCTGCGAGAAAAGTACAGAGAGCC 450  
 RESULT 7  
 US-09-864-761-16671  
 Sequence 16671, Application US/09864761  
 Patent No. US20020048763A1  
 GENERAL INFORMATION:  
 APPLICANT: Penn, Sharron G.  
 APPLICANT: Smith, David R.  
 APPLICANT: Hazenel, David K.  
 APPLICANT: Chen, Wensheng  
 TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FO  
 FILE REFERENCE: A60163-X-1  
 CURRENT FILING DATE: 2001-05-23  
 PRIOR FILING DATE: 2000-02-04  
 PRIOR APPLICATION NUMBER: US 60/180,312  
 PRIOR FILING DATE: 2000-05-26  
 PRIOR APPLICATION NUMBER: US 60/207,456  
 PRIOR FILING DATE: 2000-09-27  
 PRIOR APPLICATION NUMBER: US 09/632,366  
 PRIOR FILING DATE: 2000-08-03  
 PRIOR APPLICATION NUMBER: US 60/236,359  
 PRIOR FILING DATE: 2000-10-04  
 PRIOR APPLICATION NUMBER: US 60/234,687  
 PRIOR FILING DATE: 2000-09-21  
 PRIOR APPLICATION NUMBER: PCT/US01/00666

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## RESULT 9





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? OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 2.6
? OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 3.6
? OTHER INFORMATION: EXPRESSED IN BRCA4, SIGNAL = 3.9
? OTHER INFORMATION: EXPRESSED IN HEPATOCARCINOMA, SIGNAL = 3.4
? OTHER INFORMATION: EXPRESSED IN HL60, SIGNAL = 1.8
? OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 2.8
? OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1.8
? OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 2.3
US-09-864-761-810
Query Match
Best Local Similarity 12.5% Score 46; DB 10; Length 381;
Matches 86; Conservative 0; Mismatches 45; Indels 3; Gaps 1;

QY 170 TTGAACTGTCTTTCATCATCGTGAGCATTCCGGAGAGACTGTGAATCCAAACAGC 229
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Db 378 TGGATATCTTCGCGCTCTTCACACCTCGGCATCATGCTGAGTCAATCGCTCCAAAGAC 319
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QY 230 GCAACAATCCAGAAGACCCTTCACACAGTCAATGTA---GAGACGGCCAGGAATC 286
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 318 TGAGAGCATCTGAAGGCCCATTTCAAGCTCACATCGAGTCCGATCCCTGGCAAGAGAG 259
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QY 287 ACAAGAGCCAATC 300
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Db 258 ACAAAGCCTATGTC 245
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RESULT 15
US-09-853-386-111
? Sequence 111, Application US/09853386
? Patent No. US20020049151A1
GENERAL INFORMATION:
APPLICANT: MURPHY, Evelyn
APPLICANT: MURPHY, Evelyn
APPLICANT: COMESENLY, Orlis
APPLICANT: COMESENLY, Orlis
TITLE OF INVENTION: Therapeutic Approaches to Diseases by Suppression of the NURR1 Gene
FILE REFERENCE: P01972U51
CURRENT APPLICATION NUMBER: US/09/853,386
PRIORITY DATE: US-01-000,000
PRIORITY APPLICATION NUMBER: US-01-000,000
PRIOR FILING DATE: 2000-05-12
NUMBER OF SEQ ID NOS: 153
SOFTWARE: Patentin version 3.1
SEQ ID NO 111
LENGTH: 1146
TYPE: DNA
ORGANISM: HUMAN
US-09-853-386-111
Query Match
Best Local Similarity 9.8%; Score 36; DB 10; Length 1146;
Matches 78; Conservative 0; Mismatches 70; Indels 0; Gaps 0;

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Db 679 GGCAAGAACGCTTGAGGTGATACACATGATCTACACAGGAGCCATGATCTGCTCG 738
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QY 175 AGTGAATCTTACATCATCGTGGCATCGCTGTGAGACATGGGAAATCCAGAGACGGGA 234
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Db 739 CAGTCAATTCATATCTCTTTCATCAACATGTGTCGCATCTCATGACCAAGCTCGGSCA 798
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 235 CACTCCATGACCCCTACACCAACATGCA 262
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Db 799 TCACACAGCTGTGAGACATCTGATGATCA 826
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Job time : 138.608 secs
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GenCore version 5.1.4.p5.4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

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Title: US-09-550-163-1\_COPY\_74\_442

Product score: 369

Sequence: 1 atgctactctatccaatctt.....ctgggttaaatatgcctccc 369

Scoring table: IDENTITY-NIC

Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 112599159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 08

Maximum Match 1008

Listing first 45 summaries

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9: /SID52/gcgdata/geneseq/geneseq-emb1/NA1988.DAT.\*  
10: /SID52/gcgdata/geneseq/geneseq-emb1/NA1989.DAT.\*  
11: /SID52/gcgdata/geneseq/geneseq-emb1/NA1990.DAT.\*  
12: /SID52/gcgdata/geneseq/geneseq-emb1/NA1991.DAT.\*  
13: /SID52/gcgdata/geneseq/geneseq-emb1/NA1992.DAT.\*  
14: /SID52/gcgdata/geneseq/geneseq-emb1/NA1993.DAT.\*  
15: /SID52/gcgdata/geneseq/geneseq-emb1/NA1994.DAT.\*  
16: /SID52/gcgdata/geneseq/geneseq-emb1/NA1995.DAT.\*  
17: /SID52/gcgdata/geneseq/geneseq-emb1/NA1996.DAT.\*  
18: /SID52/gcgdata/geneseq/geneseq-emb1/NA1997.DAT.\*  
19: /SID52/gcgdata/geneseq/geneseq-emb1/NA1998.DAT.\*  
20: /SID52/gcgdata/geneseq/geneseq-emb1/NA1999.DAT.\*  
21: /SID52/gcgdata/geneseq/geneseq-emb1/NA2000.DAT.\*  
22: /SID52/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.\*  
23: /SID52/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.\*  
24: /SID52/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	369	100.0	372	22	Human #1365 for g
2	369	100.0	372	22	Human #1365 for g
3	369	100.0	372	22	Human #1365 for g
4	369	100.0	471	22	Human #1365 for g
5	369	100.0	600	22	Human #1365 for g
6	369	100.0	600	22	Human #1365 for g
7	369	100.0	655	22	Human #1365 for g
8	369	100.0	732	21	Human #1365 for g
9	369	100.0	732	24	Human #1365 for g

10	369	100.0	732	24	Human #1365 for g
11	367.4	99.6	732	21	Human #1365 for g
12	367.4	99.6	732	21	Human #1365 for g
13	367.4	99.6	732	21	Human #1365 for g
14	367.4	99.6	732	21	Human #1365 for g
15	367.4	99.6	732	21	Human #1365 for g
16	367.4	99.6	732	21	Human #1365 for g
17	367.4	99.6	732	21	Human #1365 for g
18	367.4	99.6	732	21	Human #1365 for g
19	312	84.6	312	22	Human #1365 for g
20	312	84.6	312	22	Human #1365 for g
21	312	84.6	312	22	Human #1365 for g
22	312	84.6	312	22	Human #1365 for g
23	312	84.6	312	22	Human #1365 for g
24	312	84.6	312	22	Human #1365 for g
25	312	84.6	312	22	Human #1365 for g
26	312	84.6	312	22	Human #1365 for g
27	312	84.6	312	22	Human #1365 for g
28	312	84.6	312	22	Human #1365 for g
29	312	84.6	312	22	Human #1365 for g
30	312	84.6	312	22	Human #1365 for g
31	312	84.6	312	22	Human #1365 for g
32	312	84.6	312	22	Human #1365 for g
33	312	84.6	312	22	Human #1365 for g
34	312	84.6	312	22	Human #1365 for g
35	312	84.6	312	22	Human #1365 for g
36	312	84.6	312	22	Human #1365 for g
37	312	84.6	312	22	Human #1365 for g
38	312	84.6	312	22	Human #1365 for g
39	312	84.6	312	22	Human #1365 for g
40	312	84.6	312	22	Human #1365 for g
41	312	84.6	312	22	Human #1365 for g
42	312	84.6	312	22	Human #1365 for g
43	312	84.6	312	22	Human #1365 for g
44	312	84.6	312	22	Human #1365 for g
45	312	84.6	312	22	Human #1365 for g

# ALIGNMENTS

RESULT 1	NA124432 standard; DNA; 372 BP.
1	NA124432
2	NA124432
3	NA124432
4	NA124432
5	NA124432
6	NA124432
7	NA124432
8	NA124432
9	NA124432
10	NA124432
11	NA124432
12	NA124432
13	NA124432
14	NA124432
15	NA124432
16	NA124432
17	NA124432
18	NA124432
19	NA124432
20	NA124432
21	NA124432
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23	NA124432
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28	NA124432
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35	NA124432
36	NA124432
37	NA124432
38	NA124432
39	NA124432
40	NA124432
41	NA124432
42	NA124432
43	NA124432
44	NA124432
45	NA124432

Probe #14365 for gene expression analysis in human cervical cell sample.  
Probe #14365 for gene expression, cervical epithelial cell.

Homo sapiens.

NC020015/278-A2.

09-AUG-2001.

30-JAN-2001; 2001MO-0506670.

04-FEB-2000; 2000US-0180312.

26-MAY-2000; 2000US-0207456.

30-JUN-2000; 2000US-0609408.

21-SEP-2000; 2000US-0234689.

27-SEP-2000; 2000US-0236559.

04-OCT-2000; 2000CP-0024563.

(MOLE-) MOLECULAR DYNAMICS INC.

Penn SG, Hanzel DK, Chen W, Rank DR;

XX WPI: 2001-468901/53.  
 CC Human genome-derived single exon nucleic acid probes useful for  
 PT analyzing gene expression in human cervical epithelial cells -  
 XX  
 PS Claim 25: SEQ ID No 1365: 487bp; English.  
 XX The present invention relates to human single exon nucleic acid probes  
 CC (SNPs) and the present sequence is one such probe. The SNPs are derived  
 CC from human DNA and are useful for measuring human gene expression in a  
 CC microarray which can be used for measuring human gene expression in a  
 CC sample derived from human cervical epithelial cells. By measuring gene  
 CC expression, the probes are therefore useful in grading and/or staging  
 CC of diseases of the cervix, notably cervical cancer.  
 CC Note: The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at [http://wipo.int/pub/published\\_pcl-sequences](http://wipo.int/pub/published_pcl-sequences).  
 XX  
 SQ Sequence 372 Bp; 110 A; 90 G; 82 G; 90 T; 0 other:  
 Query Match 100.0%; Score 369; DB 22; Length 372;  
 Best Local Similarity 100.0%; Pred. No. 2,4e-101;  
 Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Oy 1 ATGCTACTTTTCAATTTCACACAGCCCTGGAGACGCTTCCAGAGATTATT 60  
 Db 1 ATGCTACTTTTCAATTTCACACAGCCCTGGAGACGCTTCCAGAGATTATT 60  
 Oy 61 ACTTATATGACAAATGCGCCGACCAACAGACCTGAGACAGAGGCCCTCCAGCCAA 120  
 Db 61 ACTTATATGACAAATGCGCCGACCAACAGACCTGAGACAGAGGCCCTCCAGCCAA 120  
 Oy 121 GTTATGCTGAGAACTCTACATCTGCTGCTGACCTGAGATGAGATGAGATGTC 180  
 Db 121 GTTATGCTGAGAACTCTACATCTGCTGCTGACCTGAGATGAGATGAGATGTC 180  
 Oy 181 TCTTTCATCATCTGCGCACTCTGTGAGCACTGTGAATTCAGAGAGGAAACACTTC 240  
 Db 181 TCTTTCATCATCTGCGCACTCTGTGAGCACTGTGAATTCAGAGAGGAAACACTTC 240  
 Oy 241 AATGACCCCTACACAGACATCTTGTAGAGAGATGCGAGAAAGTACAGAGCCAAATC 300  
 Db 241 AATGACCCCTACACAGACATCTTGTAGAGAGATGCGAGAAAGTACAGAGCCAAATC 300  
 Oy 301 TTGAATCTAGAAATGCAAGAGCCACACATCTCATAGAACATGTGTGCGCTGAGTCAA 360  
 Db 301 TTGAATCTAGAAATGCAAGAGCCACACATCTCATAGAACATGTGTGCGCTGAGTCAA 360  
 Oy 361 ATGTCCCCC 369  
 Db 361 ATGTCCCCC 369  
 RESULT 2  
 AA109965  
 ID AA109965 standard; DNM: 372 Bp.  
 XX AA109965:  
 PT 09-OCT-2001 (first entry)  
 XX Probe #9356 used to measure gene expression in human breast sample.  
 RM Probs: human; breast disease; breast cancer; development disorder; 89;  
 RM Inflammatory disease; proliferative breast disease; non-carcinoma tumour.  
 XX Homo sapiens.  
 XX WO200157370-A2.  
 XX 09-AUG-2001.  
 PD

PF 29-JAN-2001; 2001WO-US000651.  
 XX 04-FEB-2000; 2000US-0180312.  
 XX 26-MAY-2000; 2000US-0207456.  
 PR 30-JUN-2000; 2000US-0608408.  
 PR 03-AUG-2000; 2000US-0632366.  
 PR 21-SEP-2000; 2000US-0234687.  
 PR 27-SEP-2000; 2000US-0236359.  
 PR 04-OCT-2000; 2000GB-0024283.  
 XX (MOLE-) MOLECULAR DYNAMICS INC.  
 XX Penn SC, Hanzel DK, Chen W, Rank DR;  
 DR WPI: 2001-476286/51.  
 XX Novel single exon nucleic acid probe used to measuring gene expression  
 PF in a human breast -  
 XX  
 PS Claim 25: SEQ ID No 9956: 322bp; English.  
 XX The present invention relates to novel single exon nucleic acid probes.  
 CC The present sequence is one such probe. The probes are useful for  
 CC measuring human gene expression in a human breast sample, where the probe  
 CC hybridises at high stringency to a nucleic acid expressed in the human  
 CC breast. The probes are useful for predicting, diagnosing, grading,  
 CC particularly those diseases with polygenic aetiology. The diseases  
 CC include: breast cancer, disorders of development, inflammatory diseases  
 CC of the breast, fibrocystic changes, proliferative breast disease and  
 CC non-carcinoma tumours.  
 CC Note: The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at [http://wipo.int/pub/published\\_pcl-sequences](http://wipo.int/pub/published_pcl-sequences).  
 XX  
 SQ Sequence 372 Bp; 110 A; 90 G; 82 G; 90 T; 0 other:  
 Query Match 100.0%; Score 369; DB 22; Length 372;  
 Best Local Similarity 100.0%; Pred. No. 2,4e-101;  
 Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Oy 1 ATGCTACTTTTCAATTTCACACAGCCCTGGAGACGCTTCCAGAGATTATT 60  
 Db 1 ATGCTACTTTTCAATTTCACACAGCCCTGGAGACGCTTCCAGAGATTATT 60  
 Oy 61 ACTTATATGACAAATGCGCCGACCAACAGACCTGAGACAGAGGCCCTCCAGCCAA 120  
 Db 61 ACTTATATGACAAATGCGCCGACCAACAGACCTGAGACAGAGGCCCTCCAGCCAA 120  
 Oy 121 GTTATGCTGAGAACTCTACATCTGCTGCTGACCTGAGATGAGATGAGATGTC 180  
 Db 121 GTTATGCTGAGAACTCTACATCTGCTGCTGACCTGAGATGAGATGAGATGTC 180  
 Oy 181 TCTTTCATCATCTGCGCACTCTGTGAGCACTGTGAATTCAGAGAGGAAACACTTC 240  
 Db 181 TCTTTCATCATCTGCGCACTCTGTGAGCACTGTGAATTCAGAGAGGAAACACTTC 240  
 Oy 241 AATGACCCCTACACAGACATCTTGTAGAGAGATGCGAGAAAGTACAGAGCCAAATC 300  
 Db 241 AATGACCCCTACACAGACATCTTGTAGAGAGATGCGAGAAAGTACAGAGCCAAATC 300  
 Oy 301 TTGAATCTAGAAATGCAAGAGCCACACATCTCATAGAACATGTGTGCGCTGAGTCAA 360  
 Db 301 TTGAATCTAGAAATGCAAGAGCCACACATCTCATAGAACATGTGTGCGCTGAGTCAA 360  
 Oy 361 ATGTCCCCC 369  
 Db 361 ATGTCCCCC 369  
 RESULT 3  
 AA500245  
 ID AA500245 standard; DNM: 372 Bp.  
 XX AA500245:  
 PT 09-AUG-2001.  
 PD



AC AAS00245;  
 DT 10-MAY-2001 (first entry)  
 XX Human potassium channel regulatory protein, Mink2, DNA sequence.  
 XX  
 XX Human: Mink2; potassium channel; cardiac arrhythmias; hypertension; ds;  
 XX angina; diabetes; renal insufficiency; urinary incontinence;  
 XX irritable colon; epilepsy; cerebrovascular ischaemia; autoimmune disease.  
 OS Homo sapiens.  
 XX  
 XX Key Location/Qualifiers  
 XX CDS 1..372  
 XX /tag=a  
 XX /product="Mink2 potassium channel protein"  
 XX  
 XX W0200114403-A1.  
 XX  
 XX 01-MAR-2001.  
 XX  
 XX 18-AUG-2000; 2000MO-US2799.  
 XX  
 XX 20-AUG-1999; 99US-0379201.  
 XX  
 XX (UYCA-) UNIT CASE WESTERN RESERVE.  
 XX  
 XX Flicker E, Wible B, Brown AM;  
 XX  
 XX WPI: 2001-218424/22.  
 XX P-PSDB: AAM00215.  
 XX  
 XX Novel potassium channel gene termed Mink2 encoding potassium channel  
 XX regulatory protein, useful for screening compounds that are useful for  
 XX treating diseases caused by aberrant potassium activity -  
 XX  
 XX Claim 1; Fig 9; 39pp; English.  
 XX  
 XX The sequence represents the coding sequence of human potassium channel  
 XX regulatory protein, Mink2. Mink2 sequence is useful for producing a  
 XX potassium channel regulatory protein useful for in vitro or in vivo  
 XX testing of agonistic or antagonistic compounds that are useful for  
 XX treating diseases caused by aberrant potassium activity. The human  
 XX cardiac arrhythmias, hypertension, angina, asthma, diabetes, renal  
 XX insufficiency, urinary incontinence, irritable colon, epilepsy, renal  
 XX cerebrovascular ischaemia, and autoimmune disease.  
 XX  
 XX Sequence 372 BP; 110 A; 90 C; 82 G; 90 T; 0 other;  
 XX  
 XX Query Match 100.0%; Score 369; DB 22; Length 372;  
 XX Best Local Similarity 100.0%; Field 0.27e-101;  
 XX Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 301 TTGATCTAGAGAAATCGAAGSCCACCATTCATGAGAAATGAGTGCAGGCTTCAAA 360  
 DB 301 TTGATCTAGAGAAATCGAAGSCCACCATTCATGAGAAATGAGTGCAGGCTTCAAA 360  
 QY 361 ATGTCCCC 369  
 DB 361 ATGTCCCC 369  
 XX  
 XX RESULT 4  
 XX AAF80269  
 XX ID AAF80269 standard; DNA; 471 BP.  
 XX AC AAF80269;  
 XX DT 29-JUN-2001 (first entry)  
 XX  
 XX Nucleotide sequence of human potassium channel subunit Isk2.  
 XX  
 XX Human: potassium channel; Isk2; gene therapy; gastric motility;  
 XX gastric acid secretion; enteric/lympanic agent; myocardial infarction; ss.  
 XX  
 XX Homo sapiens.  
 XX  
 XX Key Location/Qualifiers  
 XX CDS 79..450  
 XX /tag=a  
 XX /product="potassium channel subunit Isk2"  
 XX  
 XX W0200127246-A1.  
 XX  
 XX 19-APR-2001.  
 XX  
 XX 10-OCT-2000; 2000MO-US28014.  
 XX  
 XX 12-OCT-1999; 99US-0158781.  
 XX  
 XX (MERI) MERCK & CO INC.  
 XX  
 XX Swanson RJ, Liu Y, Folander K;  
 XX  
 XX WPI: 2001-273764/28.  
 XX P-PSDB: AAB67800.  
 XX  
 XX New DNA encoding the Isk2 potassium channel subunit, useful e.g. for  
 XX detecting mutations and screening for therapeutic agents -  
 XX  
 XX Claim 3; Fig 1A; 46pp; English.  
 XX  
 XX The present sequence encodes a human potassium channel subunit,  
 XX Isk2, and is useful for producing a protein, the protein is prob-  
 XX ably used diagnostically to detect mutations in the Isk2 gene to determine  
 XX levels of mRNA expression and to isolate homologous sequences; for  
 XX recombinant expression of Isk2; in gene therapy to increase potassium  
 XX channel activity and to generate transgenic animals, as models and  
 XX for drug screening. Recombinant Isk2 is used for studying biochemical  
 XX activity of Isk2 and its role in disorders of gastric motility and  
 XX gastric acid secretion, and to raise specific antibodies. Isk2  
 XX modulators are potentially useful for treating diseases associated with  
 XX anti-arrhythmic agents for treating myocardial infarction and as  
 XX regulators of gastric acid secretion.  
 XX  
 XX Sequence 471 BP; 143 A; 110 C; 103 G; 115 T; 0 other;  
 XX  
 XX Query Match 100.0%; Score 369; DB 22; Length 471;  
 XX Best Local Similarity 100.0%; Field 0.27e-101;  
 XX Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 61 ACTTATGACGAACTTGGCCGAGAACACAGACCTGAGACAGAGCCCTCCAAACCAAA 120  
 DB 139 ACTTATGACGAACTTGGCCGAGAACACAGACCTGAGACAGAGCCCTCCAAACCAAA 198  
 OY 121 GTTATGCTGAGAACTTCACTATGATCTGATCTGATCTGATCTGATCTGATCTGATCTG 180  
 DB 139 GTTATGCTGAGAACTTCACTATGATCTGATCTGATCTGATCTGATCTGATCTGATCTG 258  
 OY 181 TCTTTCATCATCTGTCGACATCTGTCGACATCTGTCGACATCTGTCGACATCTGTCGAC 240  
 DB 259 TCTTTCATCATCTGTCGACATCTGTCGACATCTGTCGACATCTGTCGACATCTGTCGAC 318  
 OY 241 AATGACCCCTACACAGACATCTGATGAGAGACTGCGAGGAAAGTCAAGACCAATTC 300  
 DB 319 AATGACCCCTACACAGACATCTGATGAGAGACTGCGAGGAAAGTCAAGACCAATTC 378  
 OY 301 TTGATCTGAGAAATGAGAGGACGACATCTGATGAGAGAGAGAGAGAGAGAGAGAGAG 360  
 DB 379 TTGATCTGAGAAATGAGAGGACGACATCTGATGAGAGAGAGAGAGAGAGAGAGAGAG 438  
 OY 361 ATGTCTCC 369  
 DB 439 ATGTCTCC 447  
 RESULT 5  
 ABA09192 standard; cDNA: 600 BP.  
 ID ABA09192  
 XX ABA09192:  
 DT 11-JAN-2002 (first entry)  
 XX  
 DE Human M18P1 homologue-encoding cDNA, SEQ ID NO:968.  
 XX  
 KM Human: cytokine; cell proliferation; cell differentiation; growth factor;  
 KM haematopoiesis regulation; tissue growth; immunomodulator; activin;  
 KM proliferation; tumorigenesis; chemokinesis; tumour lysis; oncogenesis;  
 KM myeloid cell disorder; lymphoid cell disorder; asthma; arthritis;  
 KM chronic inflammatory condition; proliferative retinopathy;  
 KM atherosclerosis; coronary heart disease; arterial ischaemia;  
 KM bone disorder; osteoporosis; vascular growth disorder;  
 KM tissue regeneration; wound healing; infection; immune disorder;  
 KM allergic rhinitis; drug screening; gene therapy; anti-inflammatory;  
 KM cytotoxic; osteopathic; vasotropic; cardiant; vincristine; antibacterial;  
 KM antifungal; vulnerary; antilucer; ss.  
 OS Homo sapiens.  
 XX  
 XX MO200157188-42.  
 XX  
 XX  
 XX 09-AUG-2001.  
 XX  
 XX 05-FEB-2001: 2001WO-US03800.  
 XX  
 XX 03-FEB-2000: 2000US-0496914.  
 XX 27-APR-2000: 2000US-0560875.  
 XX  
 XX (HSE-) HYSEQ INC.  
 XX  
 XX Tang YF, Liu C, Dermanac RF;  
 XX  
 XX WPI: 2001-457740/49.  
 XX P-PSDB: ABB11948.  
 XX  
 XX Human proteins and DNA encoding sequences useful for preventing,  
 XX treating or ameliorating a medical condition in a mammalian subject  
 XX e.g. arthritis and cancer -  
 XX  
 XX Claim 1: Page 826, 1963pp: English.

XX Sequences ABB10981-ABB12330 represent 1350 novel human polypeptides, and  
 CC sequences ABA08225-ABA09574 represent nucleic acids encoding them. The  
 CC sequences also represent nucleic acids encoding the novel polypeptides, a  
 CC method of the invention comprising the steps of: (a) identifying the novel  
 CC nucleic acids; (b) identifying the novel polypeptides; (c) identifying the  
 CC antibodies against the polypeptides; methods of detecting the nucleotides  
 CC or polypeptides in a sample, and methods of identifying compounds which  
 CC bind to polypeptides of the invention. Although novel, many of the  
 CC polypeptides of the invention have homology to known proteins, thereby  
 CC giving an insight into their probable biological activities, and hence  
 CC potential therapeutic applications. The polypeptides of the invention may  
 CC be used in a variety of ways, including: (a) as a growth factor or cell  
 CC differentiation activator; (b) as a growth factor activator;  
 CC haematopoietic regulatory activity; tissue growth activity;  
 CC immunomodulatory activity; tumour cell proliferation or metastasis;  
 CC chemotactic or chemokinetic activity; haemostatic, thrombotic or  
 CC thrombolytic activities; receptor or ligand activities; or may be  
 CC involved in oncogenesis, cancer cell proliferation or metastasis.  
 CC The invention is useful for preventing, treating or ameliorating medical  
 CC conditions, e.g., by protein or gene therapy. Such conditions include  
 CC cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell  
 CC disorders), chronic inflammatory conditions (e.g., asthma or arthritis),  
 CC proliferative retinopathy, atherosclerosis, coronary heart disease,  
 CC arterial ischaemia, bone disorders (e.g., osteoporosis), rheumatoid  
 CC arthritis, bone marrow failure, bone marrow transplantation, wound  
 CC repair (or nucleic acids encoding them) may be used to promote wound  
 CC healing (e.g., of burns, incisions and ulcers), while those with  
 CC immunomodulatory activities may be used in the treatment of viral,  
 CC bacterial and fungal infections in addition to immune disorders.  
 CC Polypeptides with growth factor activity may be used in cell cultures to  
 CC promote cell growth. For example, such polypeptides may be used in  
 CC the treatment of cancer, and in the treatment of diseases of the cells  
 CC that can be used to augment or replace cells damaged by illness.  
 CC autoimmune disease or accidental damage. The polypeptides and nucleotides  
 CC may also be used in the diagnosis of the above conditions, and in drug  
 CC screening techniques. The present sequence represents a cDNA encoding a  
 CC novel human polypeptide of the invention.  
 XX  
 XX Sequence 600 BP: 187 A: 133 C: 130 G: 144 T: 6 other:  
 XX  
 XX Query Match 100.0%; Score 369; DB 22; Length 600;  
 XX Best Local Similarity 100.0%; Pred. No. 3e-101;  
 XX Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 OY 1 ATGTCTCTTATTCATTTCCATTCGACAGCTGTGAGAGAGCTCTTCGAGAGATTTTAT 60  
 DB 38 ATGTCTCTTATTCATTTCCATTCGACAGCTGTGAGAGAGCTCTTCGAGAGATTTTAT 97  
 OY 61 ACTTATGACGAACTTGGCCGAGAACACAGACCTGAGACAGAGCCCTCCAAACCAAA 120  
 DB 98 ACTTATGACGAACTTGGCCGAGAACACAGACCTGAGACAGAGCCCTCCAAACCAAA 157  
 OY 121 GTTATGCTGAGAACTTCACTATGATCTGATCTGATCTGATCTGATCTGATCTGATCTG 180  
 DB 139 GTTATGCTGAGAACTTCACTATGATCTGATCTGATCTGATCTGATCTGATCTGATCTG 258  
 OY 181 TCTTTCATCATCTGTCGACATCTGTCGACATCTGTCGACATCTGTCGACATCTGTCGAC 240  
 DB 259 TCTTTCATCATCTGTCGACATCTGTCGACATCTGTCGACATCTGTCGACATCTGTCGAC 318  
 OY 241 AATGACCCCTACACAGACATCTGATGAGAGACTGCGAGGAAAGTCAAGACCAATTC 300  
 DB 319 AATGACCCCTACACAGACATCTGATGAGAGACTGCGAGGAAAGTCAAGACCAATTC 378  
 OY 301 TTGATCTGAGAAATGAGAGGACGACATCTGATGAGAGAGAGAGAGAGAGAGAGAGAG 360  
 DB 379 TTGATCTGAGAAATGAGAGGACGACATCTGATGAGAGAGAGAGAGAGAGAGAGAGAG 438  
 OY 361 ATGTCTCC 369  
 DB 439 ATGTCTCC 447

	RESULT 6
ID	AAM52645
XX	AAM52645 standard; cDNA: 600 BP.
XX	
XX	AAM52645;
DT	06-NOV-2001 (first entry)
DE	Human polynucleotide SEQ ID NO 2174.
KW	Human; cytokine; cell proliferation; cell differentiation; gene therapy; vaccine; peptide therapy; stem cell growth factor; haematopoiesis; tissue growth factor; immunomodulatory; cancer; leukemia; nervous system disorder; arthritis; inflammation; ss.
OS	Homo sapiens.
PN	MO200157190-A2.
PD	09-AUG-2001.
XX	05-FEB-2001; 2001MO-USQ4098.
PR	03-FEB-2000; 2000US-0456914.
PR	27-APR-2000; 2000US-0560875.
PR	20-JUN-2000; 2000US-0598075.
PR	19-JUL-2000; 2000US-0620325.
PR	01-SEP-2000; 2000US-0654936.
PR	15-SEP-2000; 2000US-0653561.
PR	20-OCT-2000; 2000US-0653325.
XX	30-NOV-2000; 2000US-0728422.
PA	(HYSE-) HYSEQ INC.
PI	Tang Y <sup>T</sup> , Liu C, <sup>†</sup> Dmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Mo Y; Zhao Q <sup>A</sup> , Wang D, Wang J, Zhang J, Ren F, Chen R, Wang X; Xue M <sup>J</sup> , Yang Y, Wejhtman T, Goodrich R; NPI. 2001-476283/51. P-PDB: AAM79512.
PT	Nucleic acids encoding polypeptides with cytokine-like activities,
PS	useful in diagnosis and gene therapy -
XX	Claim 1: Page 4539-4540; 6221pp; English.
XX	The invention relates to polynucleotides (AAM5156-AAM53435) and the encoded polypeptides (AAM78323-AAM80302) that exhibit activity elating to proliferation or differentiation of cell differentiation or which may induce production of other factors. The polynucleotides are useful in the vaccines or polynucleotides and polypeptides are useful in gene therapy. The vaccines or peptide therapy. The polypeptides have various cytokine-like activities, e.g., stem cell growth factor activity, haematopoiesis regulating activity, tissue growth factor activity, immunomodulatory activity and activin/inhibin activity and may be useful in the diagnosis and/or treatment of cancer, leukaemia, nervous system disorders, arthritis and other diseases.
CC	Note: Sequence for SEQ ID NO 2110 (AAM5281), 2111 (AAM5282) and 3666 (AAM80620) are omitted as the relevant pages from the sequence listing were missing at the time of publication.
SO	Sequence 600 BP; 187 A; 133 C; 130 G; 144 T; 6 other:
XX	
XX	Query Match            100.0%; Score 369; DB 22; Length 600;
XX	Best Local Similarity    100.0%; Pred. No. 3e-101;
XX	Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY	1 ATGCGACCTTATTCACAAATTTGCACAGAGCGTGGAAGAGCGTTCGAGAAGATTATTAAT 60       38 ATGCGACTTATTCACAAATTTGCACAGAGCGTGGAAGAGCGTTCGAGAAGATTATTAAT 97 
OY	61 ACTTATATGACAATATGGCCGCCAGAGAACAGGTGACCAAGGAGCCCTCAGGCAAA 120

D	b	98	ACTTATTAAGGACAAATTGGGGCGGAGAACCAACAACAGCTGCAGCAGAAGGCCGCTCCCAACGAAA	157
O	y	121	GTTATAGGTGACAAAGCTTCACATATGTGCATCCCTGACACCATATGGTAGATATGTC	180
D	b	158	TCCTAATGCTGAACACACCTACCTGCTGACACCTGTGCAATGAGAACAGAGGAAACATCCG	217
O	y	181	TCCTAATGCTGAACACACCTACCTGCTGACACCTGTGCAATGAGAACAGAGGAAACATCCG	240
D	b	218	TCTTATCATCTGTTGGGCACTCTGGTGAGCACTGTCAAATCCCAAGACGGGAAACACTCC	277
O	y	241	AATGACCCCTACACACACGACATATTTGTAGAGGACTGTGCAGSAAAGTACAAAGACCACATC	300
D	b	278	AATGACCCCTACACACACGACATATTTGTAGAGGACTGTGCAGSAAAGTACAAAGACCACATC	337
O	y	301	TTGAAATCTGAAGAAAGCAAGACAGCCACCAACCAAGCAAGATGTCGGTGGTCTCAA	360
D	b	338	TTGAAATCTGAAGAAAGTGAAGGCAACCAACCAAGCAAGATGTCGGTGGTCTCAA	397
O	y	361	ATGTCCCCC	369
D	b	398	ATGTCCCCC	406
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RESULT 7				
P	I	ID	AAK51661 standard; cDNA, 655 bp.	
X	X	AAK51661:		
A	C	XX		
D	T	XX		
X	E	XX		
X	E	XX		
K	M	XX	Human polynucleotide SEQ ID NO 206.	
K	M	XX	Human cytokine: cell proliferation; cell differentiation; gene therapy;	
K	M	XX	tissue growth factor; immunomodulatory; cancer; leukemia;	
K	M	XX	nervous system disorder; arthritis; inflammation; as.	
O	S	XX	Homio sapiens.	
P	N	XX	NC0200157190 -R2.	
P	D	XX		
P	D	XX	09-ANG-2001.	
P	F	XX	05-FEB-2001: 2001WO-US04098.	
X	X	XX		
P	R	XX	03-FEB-2000: 2000US-0496914.	
P	R	XX	27-APR-2000: 2000US-0560875.	
P	R	XX	20-JUN-2000: 2000US-0596075.	
P	R	XX	01-SEP-2000: 2000US-0624935.	
P	R	XX	15-SEP-2000: 2000US-0663561.	
P	R	XX	20-OCT-2000: 2000US-0693325.	
P	R	XX	30-NOV-2000: 2000US-0728422.	
P	A	XX	(HYSE-) HYSEQ INC.	
P	A	XX		
P	T	XX	Tung YF, Liu C, Dimaec RF, Auand V, Zhou P, Xu C, Cao Y, Ma Y;	
P	T	XX	Zhang QJ, Wang JD, Zhang Z, Zhao R, Ren F, Chen R, Wang ZM;	
P	I	XX	Xue AL, Yang Y, Wejberman T, Goodrich R,	
P	I	XX	WPI: 2001-476283/51.	
D	R	XX	P-PsDB: AAM78528.	
P	T	XX		
P	T	XX	Nucleic acids encoding polypeptides with cytokine-like activities,	
P	T	XX	useful in diagnosis and gene therapy -	
P	S	XX	Claim 1: Page 1024; 6221pp; English.	
C	C	XX	The invention relates to polynucleotides (AAK51456-AAK53435) and the	
C	C	XX	coded polypeptides (AAM7823-AAM8030) that exhibit activity elating to	
C	C	XX	cytokine, cell proliferation or cell differentiation or which may induce	

CC production of other cytokines in other cell populations. The  
 CC cytokines and polypeptides are useful in gene therapy, vaccines or  
 CC polypeptides and polypeptides are useful in gene therapy, vaccines or  
 CC e.g. stem cell growth factor activity, hematopoietic regulatory activities,  
 CC activity, tissue growth factor activity, immunomodulatory activity and  
 CC activin/inhibin activity and may be useful in the diagnosis and/or  
 CC treatment of cancer, leukemia, nervous system disorders, arthritis and  
 CC inflammation.  
 CC Information for SEQ ID NO 2110 (AKS2581), 2111 (AKS2582) and 3665  
 CC has been deposited with the European Patent Office under the name of  
 CC the applicant at the time of publication.

SO Sequence 655 BP; 196 A; 154 G; 146 G; 153 T; 6 other;

Query Match 100.0%; Score 369; DB 21; Length 655;

Local Similarity 100.0%; Pred. No. 3.1e-101; Mismatches 0; Indels 0; Gaps 0;

Mismatches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 ATGCTACTTATTCATTCACACAGCGCGGAGACGCTGCTCCAGATTTTATT 60

93 ATGCTACTTATTCATTCACACAGCGCGGAGACGCTGCTCCAGATTTTATT 152

61 ACTTATTCAGCATTCGGCGCCAGACACACACTGACAGACGCTCCACACCAA 120

153 ACTTATTCAGCATTCGGCGCCAGACACACACTGACAGACGCTCCACACCAA 212

121 GTTATGCTGAGAACTTCTATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 180

213 GTTATGCTGAGAACTTCTATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 272

181 TTTTATTCAGCATTCGGCGCCAGACACACACTGACAGACGCTCCACACCAA 240

273 TTTTATTCAGCATTCGGCGCCAGACACACACTGACAGACGCTCCACACCAA 332

241 AATGACCCCTACACAGATTCGTAGAGAGAGTGCAGAGAAAGTACAAAGCCAAATC 300

333 AATGACCCCTACACAGATTCGTAGAGAGAGTGCAGAGAAAGTACAAAGCCAAATC 392

301 TTGAACTCTGAAATTCAGAGCCACCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 360

333 TTGAACTCTGAAATTCAGAGCCACCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 452

361 ATGTCCCTC 369

453 ATGTCCCTC 461

RESULT 8  
 AAC64071 standard; cDNA; 732 BP.

AAC64071:

19-FEB-2001 (first entry)

Human potassium channel protein KCNE2 (MIRP1) cDNA, SEQ ID NO.1.

Human; KCNE2; MIRP1; potassium channel protein; KCNE1-related;

Mink-related; long QT syndrome; cardiac arrhythmia;

drug screening; knockout mouse; transgenic animal; ion channel disorder;

fast delayed rectifier potassium channel; anti-KCNE2 antibody;

HERG; 95.

Homo sapiens.

WO200063434-A1.

26-OCT-2000.

14-APR-2000; 2000MO-0510004.

15-APR-1999; 9905-0129404.

PA (UTAH ) UNIV UTAH RES FOUND.

PA (UTAH ) UNIV UTAH.

PA Abbott GW, Seest F, Splawski I, Keating MT, Goldstein SM;

DR WPI; 2000-672747/65.

DR P-PSDB; AAB25985.

XX Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for

XX diagnosing and treating ion channel disorders, especially long QT

XX syndrome.

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XX ABR6573;  
 AC 24-SEP-2002 (first entry)  
 XX  
 DE cDNA encoding human ether-a-go-go related interacting protein M1RP1.  
 XX  
 KW Human; human ether-a-go-go related gene; HERG; KCRI; M1RP1;  
 KW long QT syndrome; LQT; single nucleotide polymorphism; cardiac arrhythmia;  
 KW potassium channel; ss; gene.  
 XX  
 XX Homo sapiens.  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 74..445  
 FT /\*tag= a  
 FT /product= "M1RP1"  
 FT  
 XX W0200242735-A2.  
 XX  
 PD 30-MAR-2002.  
 XX  
 PF 30-OCT-2001; 2001W0-US45644.  
 XX  
 PR 30-OCT-2000; 2000US-244340P.  
 XX  
 XX (UTRA-) UNIV VANDERBILT.  
 XX  
 XX Balser JR, George AL, Roden DM;  
 PI WPI; 2002-527650/56.  
 DR P-PSDB; AA095168.  
 DR  
 PT Identifying a potassium channel activity modulator for drug design,  
 PT potassium channel activity modulator, potassium channel and related  
 PT cerebellar cDNA library (KCRI) polypeptide, and determining activity -  
 XX  
 XX Claim 17; Page 162-163; 164pp; English.  
 XX  
 CC The invention relates to identifying (M1) a compound that modulates  
 CC biological activity of a potassium channel (PC), by contacting a  
 CC compound with a structure comprising a PC polypeptide and a polypeptide  
 CC oligonucleotide, wherein the compound modulates the biological  
 CC activity of the PC polypeptide in the presence and absence of the  
 CC compound, where a difference in the activities indicates modulation of  
 CC biological activity of PC. Also include are identifying (M2) a candidate  
 CC compound that modulates the biological activity of a complex comprising a  
 CC human ether-a-go-go-related gene (HERG) channel polypeptide and a KCRI  
 CC polypeptide, identifying (M3) a candidate compound as a modulator of KCRI  
 CC activity, and identifying (M4) a function in a subject, comprising of a  
 CC agent, consisting of the subject, for screening (M5) for susceptibility to a drug  
 CC PC function is desired, screening (M5) for susceptibility to a drug  
 CC PC function is desired, screening (M5) for susceptibility to a drug  
 CC Induced cardiac arrhythmia in a subject, comprising obtaining a  
 CC biological sample from the subject and detecting a polymorphism of a KCRI  
 CC gene in the biological sample from the subject, where the presence of the  
 CC polymorphism indicates the susceptibility of the subject to a  
 CC drug. Also include are identifying (M6) an oligonucleotide pair, where a first  
 CC oligonucleotide of the pair hybridizes to a second portion of the KCRI  
 CC which includes a polymorphism of the KCRI gene, and the second  
 CC oligonucleotide of the pair hybridizes to a second portion of the KCRI  
 CC gene that is adjacent to the first portion and a set of antisense  
 CC oligonucleotide primers, suitable for amplifying a portion of a KCRI gene  
 CC which includes a polymorphism of the KCRI gene. (M1) is useful for  
 CC identifying a compound that modulates biological activity of PC,  
 CC identifying a function, or modulating PC function, the modulating HERG  
 CC activity in a subject, or modulating PC function, the modulating HERG  
 CC activity in a subject, or modulating PC function, the modulating HERG  
 CC compound and administering the composition, the compound is useful for  
 CC treating or preventing long QT syndrome (LQT) and is useful in drug  
 CC designing. The present sequence encodes a HERG interacting  
 CC protein M1RP1 (not defined).  
 CC  
 XX Sequence 732 BP; 221 A; 153 C; 157 G; 201 T; 0 other;

Query Match 100.0%; Score 369; DB 24; Length 732;  
 Best Local Similarity 100.0%; Pred. No. 3, 3e-101;  
 Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 ATGTCTACTTATTCACATTTCACACAGAGCGTGGAAAGAGCTTCCAGAGATTTTATTT 60  
 DB 74 ATGTCTACTTATTCACATTTCACACAGAGCGTGGAAAGAGCTTCCAGAGATTTTATTT 133  
 QY 61 ACTTATATGACAAATTTGGCCGACACACACAGAGTGCATTAAGAGCTTCCAGACAA 120  
 DB 134 ACTTATATGACAAATTTGGCCGACACACACAGAGTGCATTAAGAGCTTCCAGACAA 193  
 QY 121 GTTGACCTGTAGAAACCTGTGACATGTGCATCTGTGACATGTGACATGTGACAT 180  
 DB 194 GTTGACCTGTAGAAACCTGTGACATGTGCATCTGTGACATGTGACATGTGACAT 253  
 QY 181 TCTTTCATCATGTGCGCATCTCTGTGACACCTGTGACATCTGTAATCCAAAGACACCTCC 240  
 DB 254 TCTTTCATCATGTGCGCATCTCTGTGACACCTGTGACACCTGTGACATCTGTAATCCAA 313  
 QY 241 AATGACCCCTACACACAGTATGTGAGAGTGTGAGAGTGTGAGAGTGTGAGAGTGTG 300  
 DB 314 AATGACCCCTACACACAGTATGTGAGAGTGTGAGAGTGTGAGAGTGTGAGAGTGTG 373  
 QY 301 TTGATCTAGAGAAATGACAGCCATCATGATGAGAAATTTGGTGGCTGGTGTAA 360  
 DB 374 TTGATCTAGAGAAATGACAGCCATCATGATGAGAAATTTGGTGGCTGGTGTAA 433  
 QY 361 ATGTGCCCC 369  
 QY 434 ATGTGCCCC 442  
 DB  
 RESULT 10  
 ID AAD35170  
 XX AAD35170 standard; DNA; 732 BP.  
 XX  
 AC AAD35170;  
 XX  
 DT 25-JUL-2002 (first entry)  
 XX  
 XX Human KCNE2 wild type DNA.  
 XX  
 KW Human; Min-K related ion channel protein; M1RP1; ion channel disorder;  
 KW KCNE2; long QT syndrome; LQTS; cardiac arrhythmia; gene; ds.  
 XX  
 OS Homo sapiens.  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 74..445  
 FT /\*tag= a  
 FT /product= "Human M1RP1 protein"  
 FT  
 XX W0200222875-A2.  
 XX  
 PD 21-MAR-2002.  
 XX  
 PF 11-SEP-2001; 2001W0-US8332.  
 XX  
 PR 11-SEP-2000; 2000US-231571P.  
 XX  
 XX (UYVA) UNIV YALE.  
 PA  
 XX Goldstein SAM;  
 PT WPI; 2002-36360/39.  
 DR P-PSDB; AA022095.  
 XX  
 PT Novel gene encoding Min-K related ion channel protein subunit and  
 PT polymorphisms in this gene associated with antibiotic-induced long QT  
 XX syndrome -  
 XX

xx Claim 9; Page 43; 49pp; English.

xx The present invention relates to novel KCNE2 genes encoding Min-K related  
 CC (MIRP) 1 ion channel proteins and polymorphisms in these genes that are  
 CC associated with ion channel disorders including arrhythmic-induced long  
 CC QT syndrome (LQTS). Detecting a mutation at amino acid positions 8, 54,  
 CC 57 or 116 of MIRP1 polypeptide or a mutation at a nucleotide position  
 CC encoding the amino acid positions is useful for diagnosing the presence  
 CC of a mutation in the coding region of a KCNE2 gene.  
 CC The present invention also provides a method for diagnosing the presence  
 CC of a mutation in the coding region of a KCNE2 gene by detecting a mutation  
 CC target one or more KCNE2 polymorphisms that are associated with cardiac  
 CC arrhythmias. The present sequence is human KCNE2 wild type DNA.

xx Sequence 732 BP; 221 A; 152 C; 157 G; 202 T; 0 other;

xx Query Match 100.0%; Score 369; DB 24; Length 732;

xx Best Local Similarity 100.0%; Pred. No. 3, 3e-101; Mismatches 0; Indels 0; Gaps 0;

xx Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

xx 1 ATGCTTACTTATTCATTCACACAGCCCTGAGAGAGCTGCTCCAGAGATTTTAAAT 60

xx 74 ACTTATCTTATTCATTCACACAGCCCTGAGAGAGCTGCTCCAGAGATTTTAAAT 133

xx 61 ACTTATCTTATTCATTCACACAGCCCTGAGAGAGCTGCTCCAGAGATTTTAAAT 120

xx 134 ACTTATCTTATTCATTCACACAGCCCTGAGAGAGCTGCTCCAGAGATTTTAAAT 193

xx 121 GTTATCTGAGAGAGCTGCTCCAGAGAGCTGCTCCAGAGATTTTAAAT 180

xx 194 GTTATCTGAGAGAGCTGCTCCAGAGAGCTGCTCCAGAGATTTTAAAT 253

xx 181 TCTTATCTGAGAGAGCTGCTCCAGAGAGCTGCTCCAGAGATTTTAAAT 240

xx 254 TCTTATCTGAGAGAGCTGCTCCAGAGAGCTGCTCCAGAGATTTTAAAT 313

xx 241 AATGACCTTATTCATTCACACAGCCCTGAGAGAGCTGCTCCAGAGATTTTAAAT 300

xx 314 AATGACCTTATTCATTCACACAGCCCTGAGAGAGCTGCTCCAGAGATTTTAAAT 373

xx 301 TGAATCTGAGAGAGCTGCTCCAGAGAGCTGCTCCAGAGATTTTAAAT 360

xx 374 TGAATCTGAGAGAGCTGCTCCAGAGAGCTGCTCCAGAGATTTTAAAT 433

xx 361 ATGCTCCCTC 369

xx 434 ATGCTCCCTC 442

xx RESULT 11

xx AAC64083

xx AAC64083 standard; DNA; 732 BP.

xx MAC64083:

xx 19-FEB-2001 (first entry)

xx Human potassium channel protein KCNE2 (MIRP1) O9E mutant DNA.

xx Human: KCNE2; MIRP1; potassium channel protein; KCNE1-related;

xx Min-K related; long QT syndrome; cardiac arrhythmia; ion channel disorder;

xx drug screening; knockout mouse; fast delayed rectifier potassium channel; anti-KCNE2 antibody;

xx HERG; mutant; ds.

xx Homo sapiens.

xx Synthetic.

xx W0200063434-A1.

xx 26-OCT-2000.

xx 14-APR-2000; 2000WO-US10004.

PR 15-APR-1999; 9905-0129404.

xx (UTAH) UNIV UTAH RES FOUND.

xx (UTAH) UNIV UTAH.

xx Abort GW, Sestil F, Splawski I, Keating MT, Goldstein SAN;

xx WPI; 2000-672747/65.

xx P-PSDB; AAB29596.

xx Novel, nucleic acids encoding ion channel disorders, especially long QT

xx syndrome.

xx Claim 56; Page -; 132pp; English.

xx The invention relates to novel ion channel proteins related to

xx KCNE2 (MIRP) and to nucleic acids encoding these proteins.

xx The invention and to nucleic acids encoding these proteins.

xx respectively; human and mouse KCNE2 (MIRP1; AAB29585 and AAB29586,

xx respectively); and human and mouse KCNE4 (MIRP2; AAB29589 and AAB29590,

xx respectively). The cDNAs encoding these proteins are given in AAC64071-

xx AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier

xx potassium channels (I<sub>Kr</sub>), mutations in which are associated with long QT

xx syndrome (LQTS). The present invention provides a method for diagnosing the

xx presence of a mutation in the coding region of a KCNE2 gene by detecting a

xx mutation in the coding region of a KCNE2 gene.

xx nonhuman animals comprising a heterologous ion channel protein gene

xx of the invention, a transgenic animal comprising human KCNE2 and HERG

xx DNA, and methods of and screening drugs for treating long QT syndrome

xx using KCNE2 proteins (including mutants), nucleic acids encoding them

xx and antibodies against KCNE2 proteins. The methods, antibodies, nucleic

xx acids and antibodies are useful for diagnosing the presence of a mutation

xx in the coding region of a KCNE2 gene.

xx KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.

xx The present sequence represents DNA encoding a mutant human KCNE2

xx (MIRP1) specifically claimed for use in diagnostic and drug screening

xx methods of the invention.

xx Note: The present sequence is not shown in the specification, but is

xx derived from the wild-type human KCNE2 cDNA sequence shown on page

xx 118-119.

xx Sequence 732 BP; 221 A; 151 C; 158 G; 202 T; 0 other;

xx Query Match 99.6%; Score 367.4; DB 21; Length 732;

xx Best Local Similarity 99.7%; Pred. No. 9, 9e-101; Mismatches 1; Indels 0; Gaps 0;

xx Matches 369; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

xx 1 ATGCTTACTTATTCATTCACACAGCCCTGAGAGAGCTGCTCCAGAGATTTTAAAT 60

xx 74 ATGCTTACTTATTCATTCACACAGCCCTGAGAGAGCTGCTCCAGAGATTTTAAAT 133

xx 61 ACTTATCTTATTCATTCACACAGCCCTGAGAGAGCTGCTCCAGAGATTTTAAAT 120

xx 134 ACTTATCTTATTCATTCACACAGCCCTGAGAGAGCTGCTCCAGAGATTTTAAAT 193

xx 121 GTTATCTGAGAGAGCTGCTCCAGAGAGCTGCTCCAGAGATTTTAAAT 180

xx 194 GTTATCTGAGAGAGCTGCTCCAGAGAGCTGCTCCAGAGATTTTAAAT 253

xx 181 TCTTATCTGAGAGAGCTGCTCCAGAGAGCTGCTCCAGAGATTTTAAAT 240

xx 254 TCTTATCTGAGAGAGCTGCTCCAGAGAGCTGCTCCAGAGATTTTAAAT 313

xx 241 AATGACCTTATTCATTCACACAGCCCTGAGAGAGCTGCTCCAGAGATTTTAAAT 300

xx 314 AATGACCTTATTCATTCACACAGCCCTGAGAGAGCTGCTCCAGAGATTTTAAAT 373

xx 301 TGAATCTGAGAGAGCTGCTCCAGAGAGCTGCTCCAGAGATTTTAAAT 360

xx 374 TGAATCTGAGAGAGCTGCTCCAGAGAGCTGCTCCAGAGATTTTAAAT 433

xx 361 ATGCTCCCTC 369

Db 434 ATGTCCCC 442

RESULT 12  
AAC64084  
ID AAC64084 standard; DNA; 732 BP.

DT 19-FEB-2001 (first entry)

Human potassium channel KCNE2 (MiRP1) M54T mutant DNA

KM human; KCNE2; M182L potassium channel protein; KCCE1-related;  
 KM MinK-related; long QT syndrome; cardiac arrhythmia;  
 KM drug screening; knockout mouse; transgenic animal; ion channel disorder;  
 KM fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
 KM HERG; mutant; ds.

05	Homo sapiens.
05	Synthetic.

PN WO200063434-A1

PD 26-OCT-2000.

PF 14-APR-2000; 2000WO-US10004.

PR 15-APR-1999; 9905-0129404.

PA (UTAH) UNIV UTAH RES FOUND.  
PA (UYVA) UNIV YALE.

PI Abbott GW, Sesti F, Splawski I, Keating MT, Goldstein SAN, XY

DR WP1: 2000-672747/65  
DR P-PSDB: AAB29594

Novel nucleic acids

PT syndrome - diagnosing and creating ion channel disorders, especially long QT

PS Claim 56; Page ~; 132pp; English

The invention relates to novel ion channel proteins relating to KCNE1 (Mink) and to nucleic acids encoding them. The proteins of KCNE1 (Mink) are human and rat KCNE2 (MIRP1, AA825358 and AA823586, respectively), human and rat KCNE3 (MIRP2, AA825359 and AA823587, respectively), and human and mouse KCNE4 (AA825360 and AA823588, respectively). The cDNA encoding these proteins are given in AA864071-AA864076. KCNE2, along with HERG, forms cardiac fast delayed rectifier potassium channels (I<sub>Kr</sub>), mutations in which are associated with long QT syndrome. The invention also relates to methods of diagnosing long QT syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a mutation in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic mice with an endogenous KCNE2, KCNE3 or KCNE4 gene, ion channel protein gene, and methods of using the nucleic acids encoding the ion channel protein gene. The invention also relates to methods of diagnosing long QT syndrome using KCNE2 proteins (including mutants), nucleic acids encoding them and antibodies against KCNE2 proteins. The methods, antibodies, nucleic acids, and proteins may be used for diagnosing or treating ion channel disorders, especially long QT syndrome. Transgenic animals comprising KCNE2 and HERG are useful for testing anti-long QT syndrome drugs. (MIRP1 specifically claimed for use in diagnostic and drug screening methods of the invention.)

Note: The present sequence is not shown in the specification, but is derived from the wild-type human KCNE2 cDNA sequence shown on page 118-119.

S0 Sequence 732 BP; 221 A; 153 C; 157 G; 201 T; 0 other;

Query Match	99.68;	Score 367.4;	DB 21;	Length 732;
Best Local Similarity	99.78;	Pred. No. 9,9e-101;		
Matches 368;	Conservative	0;	Mismatches 1;	Indels 0;
			Gaps	0

32y	1	ATGCTACTTTTTCACATTTCCACACAGAGCTGTGAAGACGCTCTCCAGAGATTTTAT	60
33y	74	ATGCTACTTTTTCACATTTCCACACAGAGCTGTGAAGACGCTCTCCAGAGATTTTAT	120
34y	61	ACTTATATGATTTTGGCTGGCGACACACACAGCTGACAAAGAGCTCTCAACGCTAA	120
35y	134	ACTTATATGATTTTGGCTGGCGACACACACAGCTGACAAAGAGCTCTCAACGCTAA	192
36y	121	GTGATAGCTGAGAACTCTCTACTATGTCATCCCTGCTACTAGTGTGAGTGAATGTC	180
37y	194	GTGATAGCTGAGAACTCTCTACTATGTCATCCCTGCTACTAGTGTGAGTGAATGTC	252
38y	181	TCTTTGATCATCTGTCGCGCATCTGGTGAACCTGTGAATTCACAGAGCGGAACTCTC	240
39y	254	TCTTTGATCATCTGTCGCGCATCTGGTGAACCTGTGAATTCACAGAGCGGAACTCTC	312
40y	241	AATGACCCCTACACACATCTACTATGTGAGAGATCTGGACAGAAAATCTCAAGAGCTCAAT	300
41y	301	AATGACCCCTACACACATCTACTATGTGAGAGATCTGGACAGAAAATCTCAAGAGCTCAAT	372
42y	314	TTGAAATCTGAAAGAAATCTGAAGGCTCCATCATGAGAACTTGTGGGTGGGCTTCAAA	432
43y	361	ATGTGCTCCG 369	
44y	434	ATGTGCTCCG 442	

RESULT 13  
AAC64085  
AAC64085 standard; DNA; 732 BP.

AAC64085;

19-FEB-2001 (first entry)

Human potassium channel protein KCNE2 (MiRP1) I57T mutant DNA

Human; KCNE2; MiRP1; potassium channel protein, KCNE1-related; MiRP-related; long QT syndrome; cardiac arrhythmia;

fast delayed

Homo sapiens

syncretic.

MOZ-4C4C00070M

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PA (UYYA ) UNIV YALE.

Abbott GW, Sestl E,

WPI; 2000-672747/65...

Novel nucleic acids encoding MiRP1, MiRP2 and MiRP3, useful for diagnosing and treating ion channel disorders, especially long QT

PS Claim 56; Page -; 132pp; English.





QY 181 TCTTTCATCATCTGGGCGCATCTCTGTGAGACACTGTGAAATTCAGAGAGCGGAGACACTCC 240  
 DB 254 TCTTTCATCATCTGGGCGCATCTCTGTGAGACACTGTGAAATTCAGAGAGCGGAGACACTCC 313  
 OY 241 AATGACCCCTACACACACAGTACATTTGTAGAGACTGCGAGAAAGTACAGAGCCAAATC 300  
 DB 314 AATGACCCCTACACACAGTACATTTGTAGAGACTGCGAGAAAGTACAGAGCCAAATC 373  
 OY 301 TTGTATATGAGAAATCGAATCGACACCATTCATCATGAGAAACATTTGTGCGCGCTGCTCAA 360  
 DB 374 TTGTATATGAGAAATCGAATCGACACCATTCATCATGAGAAACATTTGTGCGCGCTGCTCAA 433  
 OY 361 ATGTGCCCC 369  
 DB 434 ATGTGCCCC 442  
 RESULT 15  
 AAD35169  
 ID AAD35169 standard; DNA; 732 BP.  
 AC AAD35169;  
 XX 25-JUL-2002 (first entry)  
 XX Human KCHN2 mutant DNA (C420T).  
 XX Human; Min-K related ion channel protein; MIRP1; ion channel disorder;  
 KM KCHN2; long QT syndrome; LQTS; cardiac arrhythmia; mutant; gene; SNP;  
 KW single nucleotide polymorphism; ds.  
 XX Homo sapiens.  
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 FT CDS 74..445  
 FT /lag- a  
 FT /product- "Human MIRP1 mutant protein"  
 FT /replace (420, C)  
 FT /tag- b  
 FT /standard\_name- "single nucleotide polymorphism (SNP)"  
 PN NO200222875-A2.  
 PD 21-MAR-2002.  
 XX 11-SEP-2001; 2001KW-US28332.  
 XX 11-SEP-2000; 2000US-231571P.  
 XX (UYTA ) UNIV YALE.  
 XX Goldstein SAN;  
 XX WPI: 2002-362360/39.  
 DR P-PSDB; AAE22094.  
 XX Novel gene encoding Min-K related ion channel protein subunit and  
 PT polymorphisms in this gene associated with antibiotic-induced long QT  
 syndrome -  
 XX  
 PS Claim 1: Page 41-42; 49pp; English.  
 XX  
 CC The present invention relates to novel KCHN2 genes encoding Min-K related  
 CC (MIRP) 1 ion channel proteins and polymorphisms in these genes that are  
 CC associated with long QT syndrome disorders including antibiotic-induced long  
 CC QT syndrome (LQTS). Detecting polymorphisms in this gene, including amino  
 CC 57 or 116 of MIRP1 polypeptide or a mutation at a nucleotide position  
 CC encoding the amino acid positions is useful for diagnosing the presence  
 CC of a polymorphism that causes drug-induced LQTS. The diagnostic methods  
 CC are useful in the development of new drug therapies which selectively  
 CC target one or more KCHN2 polymorphisms that are associated with cardiac  
 CC arrhythmias. The present sequence is human KCHN2 mutant DNA (C420T).

XX  
 SQ Sequence 732 BP; 221 A; 151 C; 157 G; 203 T; 0 other;  
 Query Match 99.6%; Score 367.4; DB 24; Length 732;  
 Description: similarity 39.7%; Fred. No. 3e-101;  
 Matches 368; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 OY 1 ATGTGACCTTATTCACAAATTTCCACAGAGAGCGTGGAAAGCTCTCCGAGAGATTTTAT 60  
 DB 74 ATGTGACCTTATTCACAAATTTCCACAGAGAGCGTGGAAAGCTCTCCGAGAGATTTTAT 133  
 OY 61 ACTTATATGAGAAATTTGGCGCTGAGACACACACAGCTGAGCAAGAGCCCTCCAGCCAAA 120  
 DB 134 ACTTATATGAGAAATTTGGCGCTGAGACACACACAGCTGAGCAAGAGCCCTCCAGCCAAA 193  
 OY 121 GTTATGCTGAGAACTTCTACTAGTACATCTGCTGACCTGATGATGATGAAATGTC 180  
 DB 194 GTTATGCTGAGAACTTCTACTAGTACATCTGCTGACCTGATGATGATGAAATGTC 253  
 OY 181 TCTTTCATCATCTGGGCGCATCTCTGTGAGACACTGTGAAATTCAGAGAGCGGAGACACTCC 240  
 DB 254 TCTTTCATCATCTGGGCGCATCTCTGTGAGACACTGTGAAATTCAGAGAGCGGAGACACTCC 313  
 OY 241 AATGACCCCTACACACAGTACATTTGTAGAGACTGCGAGAAAGTACAGAGCCAAATC 300  
 DB 314 AATGACCCCTACACACAGTACATTTGTAGAGACTGCGAGAAAGTACAGAGCCAAATC 373  
 OY 301 TTGTATATGAGAAATCGAATCGACACCATTCATCATGAGAAACATTTGTGCGCGCTGCTCAA 360  
 DB 374 TTGTATATGAGAAATCGAATCGACACCATTCATCATGAGAAACATTTGTGCGCGCTGCTCAA 433  
 OY 361 ATGTGCCCC 369  
 DB 434 ATGTGCCCC 442  
 Search completed: May 21, 2003, 20:27:32  
 Job time : 95.8202 secs



GenCore version 5.1.4.p5.4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 19:44:14 ; Search time 803.354 Seconds

(without alignments)  
13367.622 Million cell updates/sec

Title: US-09-550-163-1\_COPY\_74\_442

Perfect score: 369  
Sequence: 1 atgcctactatccatctt.....ctgggttcacaaatgcctcc 369

Scoring table: IDENTITY\_NMC  
Gapop 10.0 ; Gapext 1.0

Searched: 205640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing filter 45 summaries

Database :  
1: gb\_da:\*  
2: gb\_hlg:\*  
3: gb\_in:\*  
4: gb\_cm:\*  
5: gb\_ov:\*  
6: gb\_ph:\*  
7: gb\_pl:\*  
8: gb\_pr:\*  
9: gb\_ro:\*  
10: gb\_ro:\*  
11: gb\_ro:\*  
12: gb\_sy:\*  
13: gb\_sy:\*  
14: gb\_vl:\*  
15: gb\_vl:\*  
16: em\_ba:\*  
17: em\_fun:\*  
18: em\_hum:\*  
19: em\_hum:\*  
20: em\_mu:\*  
21: em\_mu:\*  
22: em\_ov:\*  
23: em\_pat:\*  
24: em\_ph:\*  
25: em\_pl:\*  
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36: em\_ro:\*  
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39: em\_ro:\*  
40: em\_ro:\*  
41: em\_ro:\*

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	369	100.0	732	6 AX406941	AX406941 Sequence
2	369	100.0	732	9 AF071002	AF071002 Homo sapi
3	369	100.0	809	9 AF302095	AF302095 Homo sapi
4	369	100.0	2408	9 AF000320	AF000320 Homo sapi
5	369	100.0	12000	9 AF000167	AF000167 Homo sapi
6	369	100.0	10000	9 AF000167	AF000167 Homo sapi
7	369	100.0	10000	17 AF000120	AF000120 Homo sapi
8	369	100.0	34000	9 AF001719	AF001719 Homo sapi
9	367.4	99.6	732	6 AX406943	AX406943 Sequence
10	367.4	99.6	732	6 AX406943	AX406943 Sequence
11	367.4	99.6	732	6 AX406943	AX406943 Sequence
12	367.4	99.6	732	6 AX406943	AX406943 Sequence
13	367.4	99.6	732	6 AX406943	AX406943 Sequence
14	263.4	71.8	163	10 AY050513	AY050513 Chy1 mouse
15	263.4	71.8	163	10 AY050513	AY050513 Chy1 mouse
16	261.8	70.9	14709	2 AC117904	AC117904 Rattus no
17	186.6	50.6	225	4 AF329636	AF329636 Oryctolagus
18	186.2	50.5	215	4 AF387764	AF387764 Equus cab
19	178.2	48.3	228	4 AF387764	AF387764 Equus cab
20	36	15.2	534	4 AF387764	AF387764 Equus cab
21	36	15.2	534	4 AF387764	AF387764 Equus cab
22	53.2	14.4	390	6 AF050512	AF050512 Chy1 por
23	53.2	14.4	390	6 AF050512	AF050512 Chy1 por
24	53.2	14.4	398	6 AF050512	AF050512 Chy1 por
25	53.2	14.4	402	9 H001584	H001584 Homo sapien
26	53.2	14.4	402	9 H001584	H001584 Homo sapien
27	53.2	14.4	436	9 H001584	H001584 Homo sapien
28	53.2	14.4	436	9 H001584	H001584 Homo sapien
29	53.2	14.4	436	9 H001584	H001584 Homo sapien
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31	53.2	14.4	436	9 H001584	H001584 Homo sapien
32	53.2	14.4	436	9 H001584	H001584 Homo sapien
33	53.2	14.4	436	9 H001584	H001584 Homo sapien
34	53.2	14.4	436	9 H001584	H001584 Homo sapien
35	53.2	14.4	436	9 H001584	H001584 Homo sapien
36	53.2	14.4	436	9 H001584	H001584 Homo sapien
37	53.2	14.4	436	9 H001584	H001584 Homo sapien
38	53.2	14.4	436	9 H001584	H001584 Homo sapien
39	53.2	14.4	436	9 H001584	H001584 Homo sapien
40	53.2	14.4	436	9 H001584	H001584 Homo sapien
41	50.4	13.7	390	4 MS062404	MS062404 Homo sapi
42	44.6	12.1	422	12 SYR1000	SYR1000 Homo sapi
43	44.6	12.1	422	12 SYR1000	SYR1000 Homo sapi
44	44.6	12.1	422	12 SYR1000	SYR1000 Homo sapi
45	38.6	10.5	4881	3 D5PHE12	D5PHE12 Homo sapi

## ALIGNMENTS

Result 1	Sequence	732 bp	DNA	Linear	Pat 14-JUN-2002
AX406941	AX406941	732 bp	DNA	Linear	Pat 14-JUN-2002
LOCUS	AX406941	732 bp	DNA	Linear	Pat 14-JUN-2002
DEFINITION	Sequence 3 from Patent WO022875.				
ACCESSION	AX406941.1				
VERSION	AX406941.1				
KEYWORDS	GI:21439816				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	Human Genome Project				
AUTHORS	Goldstein, S.A.				
TITLE	Polymorphisms associated with cardiac arrhythmia				
JOURNAL	Patent: WO 022875-A 3 21-MAR-2002;				

YALE UNIVERSITY (US)  
 Location/Qualifiers  
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 /organism="Homo sapiens"  
 /db="taxon:9606"  
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 /codon\_start=1  
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 /translation="MSTLSNFTGLDEVERRITFYDMKNRONTAEQALQAVDAE  
 NRYVYVILAWVIGMSPTIVAIIVTSVSKRSHSNDFHQIYEDMRKTSQILN  
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 BASE COUNT 221 a 192 c 157 g 202 t  
 ORIGIN

Query Match 100.0%; Score 369; DB 6; Length 732;  
 Best Local Similarity 100.0%; Pred. No. 8.6e-101;  
 Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 1 ATGTCACTTTATCCATTTCACAGAGCGCTGGAGACGCTCTCCGAAAGATTTTATT 60  
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 Db 74 ATGTCACTTTATCCATTTCACAGAGCGCTGGAGACGCTCTCCGAAAGATTTTATT 133  
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 Oy 61 ACTTATATGACAAATTTGGCGCGAGAACACAGCTGAGCAAGAGCCCTCCAGCCAA 120  
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 Oy 121 GTTGATGCTGAGAACTTACATATGCTGACACCTGATCGATGATTTGGAATGTC 180  
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 Db 314 AATGACCCCTACACAGTACATTTGTAGAGAGCTGGCGAGAAAGTACAGAGCCCAATC 373  
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 Oy 301 TTGAACTCTAGAGAAATTCAGAGCGCACATCCATAGAACTTGGCGGCGCTTCAA 360  
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 Db 374 TTGAACTCTAGAGAAATTCAGAGCGCACATCCATAGAACTTGGCGGCGCTTCAA 433  
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 Oy 361 ATGTCCCC 369  
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 Db 434 ATGTCCCC 442  
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RESULT 2  
 AF071002 732 bp mRNA linear PRI 29-APR-1999  
 LOCUS Homo sapiens mink-related peptide 1 mRNA, complete cds.  
 DEFINITION AF071002  
 ACCESSION AF071002.1 GI:4704422  
 KEYWORDS  
 SOURCE Homo sapiens.  
 ORGANISM Homo sapiens.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
 REFERENCE 1 (bases 1 to 732)  
 AUTHORS Abdoel, G.W., Seal, L.F., Splawski, I., Buck, M.E., Lehmann, M.H.,  
 Makiyama, T., Weidinger, M.T. and Goldstein, S.A.  
 TITLE MIRP1 forms a K<sup>+</sup> channel subunit and is associated with  
 cardiac arrhythmia.  
 JOURNAL Cell 97 (2), 175-187 (1999)  
 MEDLINE 99235979  
 PUBMED 10219239  
 REFERENCE 2 (bases 1 to 732)  
 AUTHORS Abdoel, G.W., Seal, L.F., Buck, M.E. and Goldstein, S.A.N.  
 JOURNAL Submitted (05-JUN-1999) Section of Developmental Biology and  
 Biophysics, Department of Pediatrics and Boyer Center for Molecular

Medicine, Yale University School of Medicine, 295 Congress Avenue,  
 New Haven, CT 06536, USA  
 Location/Qualifiers  
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Query Match 100.0%; Score 369; DB 9; Length 732;  
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Y 1 ATGTCACTTTATCCATTTCACAGAGCGCTGGAGACGCTCTCCGAAAGATTTTATT 60  
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 Oy 361 ATGTCCCC 369  
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 Db 434 ATGTCCCC 442  
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RESULT 3  
 AF302095 809 bp mRNA linear PRI 14-SEP-2000  
 LOCUS Homo sapiens voltage-gated K<sup>+</sup> channel subunit MIRP1 (KCNK2) mRNA,  
 complete cds.  
 DEFINITION AF302095  
 ACCESSION AF302095.1 GI:10121887  
 KEYWORDS  
 SOURCE Homo sapiens.  
 ORGANISM Homo sapiens.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
 REFERENCE 1 (bases 1 to 809)  
 AUTHORS Domenech, A., Estivill, X. and de la Luna, S.  
 TITLE Cloning of human MIRP1 cDNA  
 JOURNAL Dispublished  
 REFERENCE 2 (bases 1 to 809)  
 AUTHORS Domenech, A., Estivill, X. and de la Luna, S.  
 JOURNAL Direct Submission



JOURNAL Published Only in Database (1998)  
 REFERENCE 2 (bases 1 to 100000)  
 AUTHORS Hattori M., Ishii K., Toyoda A., Shiba T. and Sakaki Y.  
 TITLE Direct Submission  
 JOURNAL Submitted (11-MAY-1998) Masahira Hattori, Kitasato University,  
 Department of Science, JST Sequencing Laboratory, Kitasato 1-15-1,  
 Sagami-mura 228, Japan (E-mail: hattori@engc.ims.u-tokyo.ac.jp),  
 Tel: +81-78-787732, Fax: +81-78-7879561,  
 Tel: +81-78-787732, Fax: +81-78-7879561,  
 The Institute of Science, JST Sequencing Laboratory as a JST sequencing  
 Principal Investigator: Yoshiyuki Sakaki Ph.D.  
 Phone: +81-3-5449-5622, Fax: +81-3-5449-5445,  
 sakaki@engc.ims.u-tokyo.ac.jp  
 Sdb-leader: Yedayoshi Shiba Ph.D., Masahira Hattori Ph.D. The  
 sequence is submitted by Human Genome Sequencing in ALIS project of  
 Japan Science and Technology Corporation (JST)  
 5-3, Yonbancho, Chiyoda-ku, Tokyo 102-0078 Japan  
 For further information about this sequence, including its location  
 and relationship to other sequences, please visit our sequences  
 archive Web site (<http://www.alls.tokyo.jst.go.jp/HGS/top.html>)  
 or send email to webmaster@www.alls.tokyo.jst.go.jp.

FEATURES  
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 1. 100000  
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BASE COUNT 27603 a 21934 c 22513 g 27950 t

ORIGIN

Query Match 100.0%; Score 369; DB 9; Length 100000;  
 Best Local Similarity 100.0%; Pred. No. 2.5e-100;  
 Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 1 ATGTACTTATTCATTCACAGACAGCGTGGAGAGCTCTTCGGAAGATTTTATT 60  
 Db 80218 ATGTACTTATTCATTCACAGACAGCGTGGAGAGCTCTTCGGAAGATTTTATT 60  
 Y 61 ACTTAATGAGCAATTGGCGCCAGAAACACACAGCTGACAGAGAGCCCTCCAGACCAA 120  
 Db 80278 ACTTAATGAGCAATTGGCGCCAGAAACACACAGCTGACAGAGAGCCCTCCAGACCAA 120  
 Y 121 GTTGAATCTAGAGAACTTCTACTATGTCTACTCTGACTCTCATGTTGATTTGATTTG 180  
 Db 80338 GTTGAATCTAGAGAACTTCTACTATGTCTACTCTGACTCTCATGTTGATTTGATTTG 180  
 Y 181 TCTTTATCATCTGTCGGCATCTCTGTGTAGCACTGTGAATTAATTCAGAGAGGAAATCTC 240  
 Db 80398 TCTTTATCATCTGTCGGCATCTCTGTGTAGCACTGTGAATTAATTCAGAGAGGAAATCTC 240  
 Y 241 AATGACCCCTACACAGATCTTCTGATGAGAGATCTGCGAGAAAGATNACAGACCAATC 300  
 Db 80458 AATGACCCCTACACAGATCTTCTGATGAGAGATCTGCGAGAAAGATNACAGACCAATC 300  
 Y 301 TTGAATCTAGAGAACTGAGAGCCACACATCCATGAGAAACATGTGGTGGTGTCAA 360  
 Db 80518 TTGAATCTAGAGAACTGAGAGCCACACATCCATGAGAAACATGTGGTGGTGTCAA 360  
 Y 361 ATGTCTCTCC 369  
 Db 80578 ATGTCTCTCC 80586

RESULT 6  
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 LOCUS Homo sapiens genomic DNA, chromosome 21q22.1, D21S278-AMU region,  
 DEFINITION clone B2344F14-190B8, segment 3/9, complete sequence.  
 ACCESSION AP000167.1 GI:487132  
 VERSION  
 KEYWORDS HTG.

SOURCE  
 ORGANISM Homo sapiens DNA.  
 Homo sapiens  
 Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.  
 REFERENCE 1 (bases 1 to 100000)  
 AUTHORS Hattori M., Ishii K., Toyoda A., Taylor T.D., Hong-Seog P.,  
 Fujiyama A., Yada T., Tokoki Y. and Sakaki Y.  
 TITLE Home sapiens 890,231bp genomic DNA of 21q22.1 (REGION: D21S226-AMU  
 CLONE RANGE: B2344F13-190B8)  
 JOURNAL Published Only in Database (1999)  
 REFERENCE 2 (bases 1 to 100000)  
 AUTHORS Hattori M., Ishii K., Toyoda A., Taylor T.D., Hong-Seog P.,  
 Fujiyama A., Yada T., Tokoki Y. and Sakaki Y.  
 TITLE Direct Submission  
 JOURNAL Submitted (10-MAY-1999) Masahira Hattori, The Institute of Physical  
 and Chemical Research (RIKEN), Genomic Sciences Center (GSC),  
 Kitasato Univ., 1-15-1 Kitasato, Sagami-mura, Kanagawa 228-8535,  
 Japan (E-mail: hattori@engc.ims.u-tokyo.ac.jp),  
 URL: <http://hgs.alls.tokyo.jst.go.jp/>, Tel: +81-78-787732,  
 Fax: +81-78-7879561,  
 E. coli transposon insertion: The present data does not contain E.  
 coli transposon sequences which integrated in the  
 original/previous sequences. We determined the boundary between  
 the insertion and genomic sequences experimentally. removed the  
 insertion sequences, reconstructed the present data. The sequencing  
 data was deposited in the Human Genome Sequencing Project (HGP)  
 and The Institute of Physical and Chemical Research (RIKEN).  
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ORIGIN

Query Match 100.0%; Score 369; DB 9; Length 100000;  
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 Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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RESULT 7  
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 LOCUS Homo sapiens genomic DNA, chromosome 21q22.1, D21S278-AMU region,  
 DEFINITION clone B2344F14-190B8, segment 3/9, complete sequence.  
 ACCESSION AP000120.1 GI:487132  
 VERSION  
 KEYWORDS HTG.

AC AP000120: AP000120.1  
 XX 04-MAY-1999 (rel. 59, Created)  
 XX 26-SEP-1999 (rel. 61, Last updated, Version 3)  
 XX Homo sapiens genomic DNA of 21q22.1, GAT and AML related, SICS53-F4A1  
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 XX RTG.  
 XX Homo sapiens (human)  
 XX Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia;  
 XX Eutheria; Primates; Catarrhini; Homidae; Homo.  
 XX (1)  
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 RA Hiraoka M., Yamaguchi H., Imai K., Shimada J.;  
 RA Submitted (15-APR-1999) to the EMBL/Genbank/CDN databases.  
 RL Nika Hiraoka, Japan Science and Technology Corporation (JST), Advanced  
 RL Databases Department, 5-3, Yonbancho, Chiyoda-ku, Tokyo 102-0081, Japan  
 RL (E-mail:mike@cyo.jst.go.jp, URL:http://www-alls.tokyo.jst.go.jp/;  
 RL Tel:81-3-5214-8491, Fax:81-3-5214-8470)  
 RL  
 XX (2)  
 XX Hattori M., Ishii K., Toyoda A., Shiba T., Sakaki Y.;  
 XX Homo sapiens B17.199bp genomic DNA of 21q22.1 GAT and AML region\*;  
 RT Unpublished.  
 XX This sequence is conducted by Kitasato University JST sequencing  
 CC Laboratory as a JST sequencing team.  
 CC Principal Investigator:Yoshiyuki Sakaki Ph.D.  
 CC Phone:81-3-5449-5022, Fax : 81-3-5449-5445,  
 CC Email:shiba@kitasato.ac.jp  
 CC Sub-leader: Tadayoshi Shiba Ph.D., Masahito Hattori Ph.D.  
 CC The sequence is submitted by Human Genome Sequencing in ALIS  
 CC Project of JST.  
 CC Japan Science and Technology Corporation (JST)  
 CC 5-3, Yonbancho, Chiyoda-ku, Tokyo 102-0081 Japan  
 CC location and relationship about this sequence, including its  
 CC location and relationship about this sequence, please visit our  
 CC web site: http://www-alls.tokyo.jst.go.jp/AB53/  
 CC or send email to webmaster@www-alls.tokyo.jst.go.jp  
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 DEFINITION Homo sapiens genomic DNA, chromosome 21q, section 63/105.  
 ACCESSION AP001719 AL163264 BK000005  
 VERSION AP001719.1 GI:7768719  
 KEYWORDS  
 SOURCE  
 ORGANISM  
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 Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
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 REFERENCE  
 Hattori M., Fujiyama A., Taylor T.D., Watanabe H., Yada T.,  
 Park H.S., Toyoda A., Ishii K., Tokoki Y., Choi D.K., Soeda E.,  
 Ohki M., Takagi T., Sakaki Y., Taudien S., Blechschmidt K.,  
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 Zimmermann W., Rosenblatt K., Kudo H., Shibuya K., Nakasaki K.,  
 Antonarakis S.E., Minoshima S., Shimizu N., Nordtek G.,  
 Hornischer K., Barand P., Scharfe M., Schoen O., Desario A.,  
 Reichelt J., Kauer G., Bloeker H., Ransier J., Beck A., Klages S.,  
 Hennig S., Rieselmann L., Dagand E., Wehrmeyer S., Borzym K.,  
 Gardiner K., Nizetic D., Francis F., Lehnach H., Reinhardt R. and  
 Yaspo M.L.  
 The DNA sequence of human chromosome 21  
 (bases 1 to 34,000,000)  
 Hattori M., Fujiyama A., Taylor T.D., Watanabe H., Yada T.,  
 Park H.S., Toyoda A., Ishii K., Tokoki Y., Choi D.K., Soeda E.,  
 Ohki M., Takagi T., Sakaki Y., Taudien S., Blechschmidt K.,  
 Polley A., Menzel U., Delabar J., Kump K., Lehmann R.,  
 Peterson D., Reichwald K., Rump A., Schillinghabel M., Schudy A.,  
 Zimmermann W., Rosenblatt K., Kudo H., Shibuya K., Nakasaki K.,  
 Antonarakis S.E., Minoshima S., Shimizu N., Nordtek G.,  
 Hornischer K., Barand P., Scharfe M., Schoen O., Desario A.,  
 Reichelt J., Kauer G., Bloeker H., Ransier J., Beck A., Klages S.,  
 Hennig S., Rieselmann L., Dagand E., Wehrmeyer S., Borzym K.,  
 Gardiner K., Nizetic D., Francis F., Lehnach H., Reinhardt R. and  
 Yaspo M.L.  
 Submitted (10-APR-2000) The Chromosome 21 Mapping and Sequencing  
 Consortium: \* RIKEN Genomic Sciences Center, Human Genome Research  
 Group \* Institute of Molecular Biotechnology, Genome Analysis \*  
 Keio University School of Medicine, Dept. of Molecular Biology \*  
 GCR, Dept. of Genome Analysis \* Max-Planck Institute for Molecular





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Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 361 ATGTGCCCC 369
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Db 301787 ATGTGCCCC 301795

RESULT 9
LOCUS      AX406939
DEFINITION Sequence 1 from Patent WO222875.
ACCESSION AX406939
VERSION    AX406939.1 GI:21439814
KEYWORDS
SOURCE     human.
ORGANISM   Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1
AUTHORS   Goldstein,S.A.
TITLE     Polymorphisms associated with cardiac arrhythmia
JOURNAL   Patent: WO 022875-A 1 21-MAR-2002;
            TALE UNIVERSITY (US)
FEATURES
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                        /db_xref="gi:21439815"
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                /note="The drug associated here was quinaldine."

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Best Local Similarity 99.7%; Pred. No. 3.3e-100;
Matches 368; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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DEFINITION Sequence 5 from Patent WO222875.
ACCESSION AX406943
VERSION    AX406943.1 GI:21439818
KEYWORDS
SOURCE     human.
ORGANISM   Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1
AUTHORS   Goldstein,S.A.
TITLE     Polymorphisms associated with cardiac arrhythmia
JOURNAL   Patent: WO 022875-A 3 21-MAR-2002;
            TALE UNIVERSITY (US)
FEATURES
            Location/Qualifiers
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variation
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            99.6%; Score 367.4; DB 6; Length 732;

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Best Local Similarity 99.7%; Pred. No. 2,6e-100;  
Matches 368; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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OY 1 ATGCTCACTTATTCACATTTCCACAGACGCTGGAGAACGCTCTCCGAAAGGATTTTATT 60
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Db 434 ATGTCCCCC 442

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RESULT 11

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ACCESSION AX406945  
VERSION AX406945.1 GI:21439820  
KEYWORDS  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1 Goldstein S.A. Polymorphism associated with cardiac arrhythmia  
JOURNAL PNAS 99:1022875-8 7 21-MAR-2002;  
TITLE Polymorphism associated with cardiac arrhythmia  
JOURNAL YALE UNIVERSITY (US)  
FEATURES  
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BASE COUNT 221 a 153 c 157 g 201 t  
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Best Local Similarity 99.7%; Pred. No. 2,6e-100;  
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OY 1 ATGCTCACTTATTCACATTTCCACAGACGCTGGAGAACGCTCTCCGAAAGATTTTATT 60
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RESULT 12

LOCUS AX406947 732 bp DNA linear PAT 14-JUN-2002  
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ACCESSION AX406947  
VERSION AX406947.1 GI:21439822  
KEYWORDS  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1 Goldstein S.A. Polymorphism associated with cardiac arrhythmia  
JOURNAL PNAS 99:1022875-9 9 21-MAR-2002;  
TITLE Polymorphism associated with cardiac arrhythmia  
JOURNAL YALE UNIVERSITY (US)  
FEATURES  
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Query Match 99.6%; Score 367.4; DB 6; Length 732;  
Best Local Similarity 99.7%; Pred. No. 2,6e-100;  
Matches 368; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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OY 61 ACTTATATGACAAATTTGGCGCCAGAACCAACAGCTGAGACAGAAAGGCGCTCCCAAGCCAA 120
Db 134 ACTTATATGACAAATTTGGCGCCAGAACCAACAGCTGAGACAGAAAGGCGCTCCCAAGCCAA 193

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 DB 434 ATGTCTCCCC 442  
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RESULT 13  
 BC022699  
 LOCUS Mus musculus, 1664 bp mRNA linear ROD 07-AUG-2002  
 DEFINITION RIKEN cDNA 2200002116 gene, clone MGC:31447  
 IMAGE:4481325, mRNA, complete cds.  
 ACCESSION BC022699.1 GI:18490550  
 VERSION MGC.  
 KEYWORDS  
 SOURCE house mouse.  
 ORGANISM Mus musculus.

REFERENCE  
 AUTHORS Mammalia: Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 TITLE Direct Submission  
 JOURNAL Submitted (01-FEB-2002) National Institutes of Health, Mammalian  
 Gene Collection (MGC), Cancer Genomics Office, National Cancer  
 Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,  
 USA

REMARK  
 COMMENT With MGC Project URL: http://mgc.ncl.nih.gov  
 Email: cgrapp@femail.nih.gov

Tissue Procurement: The Cepho Laboratory  
 cDNA Library Preparation: Life Technologies, Inc.  
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (ILMI),  
 DNA Sequencing by: Sequencing Group at the Stanford Human Genome  
 Center, Stanford University School of Medicine, Stanford, CA 94305  
 Gen Bank: http://www.ncbi.nlm.nih.gov/GenBank/GenBank.html  
 Gen Bank: (Diction: MGC) Stanford.edu  
 Dickson, M., Schmitt, J., Gilwood, D., Rodriguez, A., and Myers,  
 R. M.

Clone distribution: MGC clone distribution information can be found  
 through the I.M.A.G.E. Consortium/ILMI at: http://image.llnl.gov  
 Series: RIKEN Plate: 44 Row: a Column: 22  
 The following clone was selected for full length sequencing because it  
 passed the following selection criteria: Hexamer frequency OK  
 analysis

FEATURES  
 source

1..1664 Location/Qualifiers  
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 /clone\_lib="NIH\_MGC\_94"  
 /lab\_host="DH10B"  
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 BASE COUNT 502 a 537 c 404 g 421 t  
 ORIGIN

Query Match 71.8%; Score 265; DB 10; Length 1664;  
 Best Local Similarity 82.4%; Pred. No. 3,6e-69;  
 Matches 304; Conservative 0; Mismatches 65; Indels 0; Gaps 0;  
 QY 1 ACTGTCCTTATTCATATTCACACAGCCGCGAGACAGCTCTCCAAATTTTAT 60  
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 DB 90 ATGGCACAATTGACCAATTGACCCAGACACTGGAGGATGCTTCAAAAAGTTTATT 149  
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 QY 61 ACTTATATGACATTTGGCCGACAGACACACAGCTGACACAGAGCCCTCCAAACAA 120  
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 DB 150 ACTTATATGACAGCTGACAGACACACAGCTGACACAGAGCCCTCCAAACAA 189  
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 QY 121 GTGATGCTGAGAACTTCTACTAGTATCTCTGTAACCTTGTATGATGATGTC 200  
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 DB 210 GTGATGCTGAGAACTTCTACTAGTATCTCTGTAACCTTGTATGATGATGTC 269  
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 QY 181 TCTTTCATCATGTCGGCATTCCGTGGAGACATGCGAAATCCAAAGACGGGAACTCC 240  
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 DB 270 TCTTTCATCATGTCGGCATTCCGTGGAGACATGCGAAATCCAAAGACGGGAACTCC 329  
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 QY 241 AATGACCCCTACACACATGATCTGTATGAGAGACTGGACGGAATAAGTACAGGCAATC 300  
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 DB 330 CACACCCCTACACACATGATCTGTATGAGAGACTGGACGGAATAAGTACAGGCAATC 389  
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 QY 301 TTGATATCTGAGAAATCATGAGACCCATCATCAATGATGATGGTGGCTGGTCCAA 360  
 |||||  
 DB 390 TCTGATATCTGAGAAATCATGAGACCCATCATCAATGATGATGGTGGCTGGTCCAA 449  
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 QY 361 ATGTCTCCCC 369  
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 DB 450 ATGTCTCCCC 458  
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RESULT 14  
 BC022699  
 LOCUS Mus musculus, 372 bp mRNA linear ROD 15-OCT-2001  
 DEFINITION RIKEN cDNA 2200002116 gene, clone MGC:31447  
 ACCESSION AY050513.1 GI:1615156  
 VERSION AY050513.1  
 KEYWORDS  
 SOURCE Cavia porcellus.

REFERENCE  
 AUTHORS Mammalia: Eutheria; Rodentia; Sciurognathi; Caviidae; Cavia.  
 TITLE Direct Submission  
 JOURNAL Submitted (12-AUG-2001) Physiology, Virginia Commonwealth  
 University, 1101 East Marshall Street, Richmond, VA 23298, USA

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 Location/Qualifiers  
 /organism="Cavia porcellus"  
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Query Match      71.4%: Score 263.4; DB 10; Length 312;
Best Local Similarity 82.1%: Pred. No. 8-36-99;
Matches 303; Conservative 0; Mismatches 66; Indels 0; Gaps 0;

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Oy 61 ACTTATATGACAACTTATGACGACGACGACGACGACGACGACGACGACGACGAC 120
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Oy 121 GTTATGCTGAGAACTTATGACGACGACGACGACGACGACGACGACGACGACGAC 180
Db 121 GTTATGCTGAGAACTTATGACGACGACGACGACGACGACGACGACGACGACGAC 180
Oy 181 TCTTTTCATCTGCTGACGACGACGACGACGACGACGACGACGACGACGACGAC 240
Db 181 TCTTTTCATCTGCTGACGACGACGACGACGACGACGACGACGACGACGACGAC 240
Oy 241 ATGACCCCTACACAGTATGATGATGACGACGACGACGACGACGACGACGACGAC 300
Db 241 ATGACCCCTACACAGTATGATGATGACGACGACGACGACGACGACGACGACGAC 300
Oy 301 TTGAACTCTGAGAACTTATGACGACGACGACGACGACGACGACGACGACGACGAC 360
Db 301 TTGAACTCTGAGAACTTATGACGACGACGACGACGACGACGACGACGACGACGAC 360
Oy 361 ATGTCCCTC 369
Db 361 GTGTCAACC 369

RESULT 15
LOCUS AF071003 468 bp mRNA linear ROD 29-Apr-1999
DEFINITION Rattus norvegicus milk-related peptide 1 mRNA, complete cds.
ACCESSION AF071003.1 GI:4704424
VERSION AF071003.1
KEYWORDS
SOURCE Rattus norvegicus.
ORGANISM Rattus norvegicus.
Eukaryota; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
REFERENCE 1 (bases 1 to 468)
AUTHORS Abbott-G.W., Sestl.F., Splawski,I., Buck M.E., Lehmann M.H.,
Timothy K.W., Keating M.T. and Goldstein, S.A.
MIRP forms 1K potassium channels with HERG and is associated with
cardiac arrhythmia
JOURNAL Cell 97 (2), 175-187 (1999)
MEDLINE 99235979
RECORD 10219239
REFERENCE 1 (bases 1 to 468)
AUTHORS Abbott-G.W., Sestl.F., Buck M.E. and Goldstein, S.A.N.
DIRECT SUBMISSION
SUBMITTED (05-JUN-1998) Section of Developmental Biology and
Biophysics, Department of Pediatrics and Boyer Center for Molecular
Medicine, Yale University School of Medicine, 295 Congress Avenue,
New Haven, CT 06536, USA
FEATURES
SOURCE Local/Gen/Dual/alters
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BASE COUNT 118 a 126 c 111 g 93 t
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Best Local Similarity 82.1%: Pred. No. 8-36-99;
Matches 303; Conservative 0; Mismatches 66; Indels 0; Gaps 0;

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Db 61 ACTTATATGACAACTTATGACGACGACGACGACGACGACGACGACGACGACGAC 120
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Oy 121 GTTATGCTGAGAACTTATGACGACGACGACGACGACGACGACGACGACGACGAC 180
Db 121 GTTATGCTGAGAACTTATGACGACGACGACGACGACGACGACGACGACGACGAC 180
Oy 155 GTGATGCTGAGAACTTATGACGACGACGACGACGACGACGACGACGACGACGAC 214
Db 155 GTGATGCTGAGAACTTATGACGACGACGACGACGACGACGACGACGACGACGAC 214
Oy 181 TCTTTTCATCTGCTGACGACGACGACGACGACGACGACGACGACGACGACGAC 240
Db 181 TCTTTTCATCTGCTGACGACGACGACGACGACGACGACGACGACGACGACGAC 240
Oy 215 GCTTTCACTGTGTGCTGACGACGACGACGACGACGACGACGACGACGACGACGAC 274
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Oy 241 ATGACCCCTACACAGTATGATGATGACGACGACGACGACGACGACGACGACGAC 300
Db 241 ATGACCCCTACACAGTATGATGATGACGACGACGACGACGACGACGACGACGAC 300
Oy 275 CAGACCCCTACACAGTATGATGATGACGACGACGACGACGACGACGACGACGAC 334
Db 275 CAGACCCCTACACAGTATGATGATGACGACGACGACGACGACGACGACGACGAC 334
Oy 301 TTGAACTCTGAGAACTTATGACGACGACGACGACGACGACGACGACGACGACGAC 360
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Oy 361 ATGTCCCTC 369
Db 361 GTGTCAACC 403

Search completed: May 21, 2003, 21:21:35
Job time : 1122.35 secs

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GenCore version 5.1.4-p5.4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: May 15, 2003, 14:21:24 ; Search time 44 Seconds

(without alignments)  
268,739 Million cell updates/sec

Title: US-09-550-163-2

Perfect score: 632  
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Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283224 seqs, 96134422 residues

Total number of hits satisfying chosen parameters: 283224

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 08

Maximum Match 1008

Listing first 45 summaries

Database :

PIR\_73:\*  
1: pir1:\*  
2: pir2:\*  
3: pir3:\*  
4: pir4:\*

Prod. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

No.	Score	Query	Length	DB	ID	Description
1	133.5	21.0	129	1	A32447	potassium channel-
2	128.5	20.3	129	2	S17307	potassium channel-
3	128	20.3	130	2	A35633	potassium channel
4	126.5	20.0	125	2	A49392	min K potassium ch
5	126.5	20.0	125	2	148146	potassium channel-
6	98.5	13.4	439	2	D64510	hypothetical prote
7	76.5	12.1	557	2	B47700	hypothetical prote
8	76.5	12.1	557	2	B47700	hypothetical prote
9	75.5	11.9	367	2	E81065	proteasome, proba
10	75.5	11.9	3839	2	T49799	related to TOM1 pr
11	75.5	11.9	1003	2	AH2335	toxin secretion AB
12	74.5	11.8	1042	2	H70303	isolectin-B4A11
13	74.5	11.8	928	2	T41332	hypothetical prote
14	74.5	11.6	451	2	E83244	hypothetical prote
15	73.5	11.6	451	2	E83244	hypothetical prote
16	73.5	11.6	652	2	T41162	hypothetical prote
17	73	11.6	400	2	B64071	tyrosine-specific
18	73	11.6	591	2	AD2148	two-component sens
19	73	11.6	1224	2	E71611	hypothetical prote
20	72.5	11.5	371	2	D89995	accessory gene reg
21	71	11.2	174	2	AE6234	hypothetical prote
22	71	11.2	281	2	AE6051	H+/K+-exchanging A
23	70.5	11.2	431	2	F43131	hypothetical prote
24	70.5	11.2	431	2	F43131	hypothetical prote
25	70	11.1	532	2	C90073	hypothetical prote
26	70	11.1	1357	2	T16860	probable adl - Msc
27	70	11.1	1357	2	T16860	hypothetical prote
28	65.5	11.0	554	2	T27878	hypothetical prote
29	65.5	11.0	946	2	S48255	probable membrane

30	69.5	11.0	1154	2	T39663	paired amphiphatic
31	69.5	11.0	3973	2	B71612	hypothetical prote
32	69	10.9	282	1	E69906	conserved hypotet
33	69	10.9	548	2	E89910	glycine betaine tr
34	68.5	10.8	350	2	B81803	probable secreted
35	68.5	10.8	655	2	H96992	probable receptor
36	68.5	10.8	696	2	C85047	probable transposo
37	68.5	10.8	701	2	S35113	TiP1 protein - yea
38	68.5	10.8	1007	2	S48335	rho-type GTPase-ac
39	68.5	10.8	1911	2	T43048	calcium channel al
40	68	10.8	503	2	T15894	hypothetical prote
41	68	10.8	378	2	G37501	uncharacterized co
42	68	10.8	378	2	G37501	uncharacterized co
43	68	10.8	435	2	A12734	AAC transporter, m
44	68	10.8	435	2	A12734	AAC transporter, m
45	68	10.8	464	2	T20238	hypothetical prote

## ALIGNMENTS

RESULT 1  
A:32447  
M:Alternate names: delayed rectifier potassium channel  
C:Species: Homo sapiens (man)  
C:Date: 28-Jan-2000 /sequence-revision 28-Jan-2000 /text-change 21-Jul-2000  
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A:Accession: J017307  
A:Molecule type: mRNA  
R:Cross-references: DBML:X60457; NID:g952779; PIDN:CMA42900.1; PID:g952780

R:Keywords: phosphoprotein; transmembrane protein; beta-amyloid precursor protein; Alzheimer's disease; amyloid plaques; A-beta peptide; APP processing; gamma-secretase; Notch signaling; presenilin; PS1; PS2; FERM Lefc.; Cdk1; 166-172; 1992

A>Title: I-SKI, a slowly activating voltage-sensitive K(+) channel. Characterization of

A:Reference number: S21135; MWID:92233938; PMID:1568475

A:Accession: S21135  
A:Molecule type: mRNA  
A:Medline: 1-129 <L&S>  
C:Cross-references: GB:M22412; NID:g923779; PIDN:CMA42900.1; PID:g952780  
C:Superfamily: human potassium channel-activating protein  
C:Keywords: phosphoprotein; potassium channel; transmembrane protein  
F:44-66/Domains: transmembrane status predicted <TM>

Query Match 20.3%; Score 128.5; DB 2; Length 129;  
Best Local Similarity 35.0%; Pcd 1e-06;  
Matches 36; Conserved: 22; Mismatches 28; Indels 17; Gaps 6;

OY 26 MHQGTAT-----ELMAYNDKMFVITYITIMWISGSPVATYSVSRR 77  
|||::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|:  
DB 17 MOE--TADGGNNGLARRSQLRD---LENTYLILALFGFPFLTGIMSYRSKL 71  
|||||::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|:

OY 78 EHSDDHYGYVED--MQEKYS--QLMLESKAT-IHENGIA 116  
|||::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|:  
DB 72 EHSDPPNYTESDMOERKAIVFOARVLSEFRACVLENQA 114

RESULT 3

Potassium channel, protein - rat  
P:Accession: X56533  
N:Alternate names: I-R-K+ channel protein  
C:Species: Rattus norvegicus (Norway rat)  
C:Length: 225 amino acids; sequence position 1-225 Sep-1990 \*text.Change 21-Jul-2000  
C:Medline: 136363; PubMed:1515393; NID:g923779; PIDN:CMA42900.1; PID:g952780  
R:ProteinData: K.J.S.: Attnavayge.27; Bennett, C.; Stein, R.B.; Samson, R.  
Proc. Natl. Acad. Sci. U.S.A. 87, 2975-2979, 1990

A>Title: Cloning and expression of the delayed-rectifier I-R-K channel from neonatal rat

A:Reference number: A35633; MWID:90222152; PMID:2183220

A:Accession: A35633  
A>Status: preliminary  
A:Molecule type: mRNA  
A:Medline: 1-130 <CD>  
R:Cross-references: GB:M4641; NID:g203976; PIDN:AAM41098.1; PID:g203977

A>Title: Cloning of a membrane protein that induces a slow voltage-gated potassium current

A:Reference number: A33177; MWID:8905617; PMID:3194754

A:Accession: A33177  
A>Status: preliminary  
A:Molecule type: mRNA  
A:Residues: 1-130 <TR>

A:Cross-references: GB:M22412; NID:g206051; PIDN:AAM41822.1; PID:g206052

A:Prognell, H.; Shan, K.Y.; Timmer, J.S.; MacLusky, N.J.; Natholin, F.; Kaczmarek, L.K.

A>Title: Estrogen induction of a small, putative K+ channel mRNA in rat uterus.

A:Reference number: J00140; MWID:90267331; PMID:2344412

A:Accession: J00140  
A:Molecule type: mRNA  
A:Residues: 1-130 <PR>

A:Experimental source: uterus  
R:Bloom, M.; Masu, M.; Tsuchida, K.; Mori, T.; Ohkubo, H.; Nakashima, S.

J. Neurosci. 10B, 200-206, 1990

A>Title: Characterization of gene organization and generation of heterogeneous mRNA species

A:Reference number: I55193; MWID:9103347; PIDN:Z229022

A:Accession: I55193  
A>Status: partial  
A:Molecule type: DNA  
A:Sequence analysis: translated from CB/BNBL/DDBT

A:Cross-references: GB:D10708; NID:g220508; PIDN:BA01552.1; PID:g2160466

A:Superfamily: human potassium channel-activating protein  
R:Keywords: phosphoprotein; transmembrane protein

```
F: 45-67/Region : hydrophobic

Query Match
Best Local Similarity    20.3% ; Score 128 ; DB 2; Length 130;
Matches      29; Conservative   17; Mismatches       4; Gaps     3;

QY      51 LYLAWMGPSFIVALTSTVSKRRRESDNPDIHYVED-MOEKYS-QILTLEESK 107
        |||::|| | ::|||:: |||::|| | |||::|| | |||::|| |
Db       46 LYIWLDFPGFFTLGIMLTIIRSKKLHSHDPNVYTIESDMOKGALFOARLESFR 105

QY      108 AF-IHENGA 116
Db       106 ACVIENQA 115


RESULT 4
A49392
min K potassium channel, gpisk - guinea pig
C:/Species: Cavia porcellus (guinea pig)
C:/Date: 07-Apr-1992 ; Sequence: revIdon 18-Nov-1994 %text_change 10-Dec-1999
Rytanum M.D.; Busch A.E.; Bond, C.T.; Maylie, J.; Adelman, J.P.
Proc. Natl. Acad. Sci. U.S.A. 90, 11520-11532, 1993
A/File: The min K channel underlies the cardiac potassium current Iks and mediates s
A:/Reference number: A49392; MTID:94089666; PMID:8265583
A/Accession: A49392
A/status: preliminary; not compared with conceptual translation
A/molecule type: nucleic acid
A/codon usage: 11.2% GC
A/experimental source: heart
A/Note: sequence extracted from NCBI backbone (MCSIP:140983)
C:/Superfamily: human potassium channel-activating protein

Query Match
Best Local Similarity    20.0% ; Score 126.5 ; DB 2; Length 125;
Matches      22; Conservative   15; Mismatches       11; Indels   1; Gaps     1;

QY      51 LYLWMMGSPFIVALTSTVSKRRRESDNPDIHYVED-MOEKYS 98
        |||::|| | ::|||:: |||::|| | |||::|| | |||::|| |
Db       45 LYIWLDFPGFFTLGIMLTIIRSKKLHSHDPNVYTIESDMOKMDKA 93


RESULT 5
IA4816
potassium channel-activating protein - Cavia cobaya
C:/Species: Cavia cobaya
C:/Date: 04-Sep-1997 ;sequence:revIdon 04-Sep-1997 %text_change 31-Jan-2000
C/Acession: IA4816
R/hang: Y.J.; Jurkiewicz, N.K.; polander, K.; Lazarides, E.; Salata, J.J.; Swanson, R.
Proc. Natl. Acad. Sci. U.S.A. 91, 1766-1770, 1994
A/Reference number: A53135; MTID:94173910; PMID:7510407
A/Accession: IA4816
A/status: preliminary; translated from GB/EMBL/DDBJ
A/molecule type: mRNA
A/bioassays: A:125 <RS>-
A/Cross-references: GS:NID:g484140; PDB:AAA73394.1; PID:g484141
C/Superfamily: human potassium channel-activating protein

Query Match
Best Local Similarity    20.0% ; Score 126.5 ; DB 2; Length 125;
Matches      22; Conservative   15; Mismatches       11; Indels   1; Gaps     1;

QY      51 LYLWMGMSPFIVALTSTVSKRRRESDNPDIHYVED-MOEKYS 98
        |||::|| | ::|||:: |||::|| | |||::|| | |||::|| |
Db       45 LYIWLDFPGFFTLGIMLTIIRSKKLHSHDPNVYTIESDMOKMDKA 93


hypercalcaemic protein MJCELO4 - Methanococcus jannaschii plasmid pMRB800
D664510
[Species: Methanococcus jannaschii]
>Date: 13-Sep-1996 %sequence:revIdon 13-Sep-1996 %text_change 08-Oct-1999
```



[illegible]



Query Match	11.6%	Score 73.5;	DB 2;	Length 171;
Best Local Similarity	36.6%	Pred. No. 4.2;		
Matches 15;	Conservative 11;	Mismatches 12;	Indels 3;	Gaps 1

```

Qy 50 ILIWMNIGMSFT---IVALVSTYTKSKRRHSNDPHY 87
      :::::  ||  ::::  ::  ::  ::  ::  ::  ::
Db 121 VFLVTLIAAVSFIGVALMLVLLIKNPKNHSKKPMAY 161

```

## RESULT 15

A: CIGOS-references: EMBL:247547; NID:g103057; PIDN:CAAG7619.1; PID:g13334497  
A: Experimental source: female gametophytes  
A: Note: The nucleotide sequence was submitted to the EMBL Data Library, January 1995  
C: Genetics:  
I: Gene: nsa7n

**C/Superfamily:** NADH dehydrogenase (ubiquinone) chain 2

Query Match	11.68;	Score 73.5;	DB 2;	Length 497;
Best Local Similarity	26.28;	Pred. No. 13;		
Matches 16;	Conservative 14;	Mismatches 14;	Indels 17;	Gaps 2

```
QY      43 AENFYVILLMVMIGMSFIYALIVSVKSKRHSNDPYHQIYVEDWQZKKKSQLLN 102
          : : ||| : : | : ||| :           | |       : : : :
Db     335 SSSIFYVILIYITMIGIFSILLT-----RYNYSYH-----YYCRYLQDLIS 377
```

QY	103	L	103
		1	
Db	378	L	378

Search completed: May 15, 2003, 14:25:36  
Job time : 47 secs



GenCore version 5.1.3  
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 24, 2002, 13:15:00 Search time 1800 Seconds  
(without alignments)  
4324.818 Million cell updates/sec

Title: US-09-550-163-1\_COPY\_74\_445

Sequence: 372  
1 agtgcactatccatctt.....gtctcaaatgtcccccgga 372

Scoring table: IDENTITY\_NUC  
Gapop 10.0, Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues

Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :  
1: gb\_ba:\*  
2: gb\_hlg:\*  
3: gb\_in:\*  
4: gb\_cm:\*  
5: gb\_cm:\*  
6: gb\_cm:\*  
7: gb\_cm:\*  
8: gb\_cm:\*  
9: gb\_cm:\*  
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45: gb\_cm:\*

## SUMMARIES

Prod. No. is the number of results predicted by chance to have a score greater than or equal to the highest score observed printed, and is derived by analysis of the total score distribution.

Result No. Score Match Length DB ID Description

Result	No.	Score	Match Length	DB ID	Description
1	372	100.0	732	9	AF071002
2	372	100.0	809	9	AF302095
3	372	100.0	24608	9	AP000320
4	372	100.0	100000	9	AP000052
5	372	100.0	100000	9	AP000120
6	372	100.0	100000	9	AP000120
7	372	100.0	340000	9	AP001215
8	268	72.0	1664	10	BC022659
9	265.4	71.6	372	10	AF050513
10	265.4	71.6	468	10	AF071003
11	186.2	50.1	215	4	AF329636
12	178.2	47.9	228	4	AF387764
13	13	10.1	524	10	RA315812
14	56	13.1	524	10	RA315812
15	53.2	14.3	390	9	AF135188
16	53.2	14.3	390	10	AF050512
17	53.2	14.3	398	6	I40373
18	53.2	14.3	402	9	H0M1SKA
19	53.2	14.3	408	9	H0M1SKA
20	53.2	14.3	408	9	H0M1SKA
21	53.2	14.3	408	9	H0M1SKA
22	53.2	14.3	1703	6	AR164693
23	53.2	14.3	43126	9	AP000324
24	53.2	14.3	100000	9	AP000053
25	53.2	14.3	100000	9	AP000121
26	53.2	14.3	100000	9	AP000168
27	53.2	14.3	340000	9	AP001720
28	52.6	14.2	931	10	AB032575
29	52.6	14.2	931	10	AB032575
30	51	13.7	471	10	RA315812
31	51	13.7	471	10	RA315812
32	50.4	13.5	390	4	MS062404
33	44.6	12.0	422	12	SYNMINK
34	41	11.0	608	6	AX346023
35	39	10.5	16182	2	AP004643
36	38.2	10.3	16685	2	AC011594
37	38.2	10.3	17045	2	AC015970
38	38.2	10.3	179522	2	AC104231
39	37.2	10.0	346294	1	AP002999
40	37.2	9.8	2652	6	AF75965
41	36.6	9.8	2652	6	AF75965
42	36.6	9.8	2652	6	AF75965
43	36.6	9.8	2652	6	AF75965
44	36.6	9.8	2652	6	AF75965
45	36.6	9.8	2652	6	AF75965

## ALIGNMENTS

RESULT 1  
AF071002 732 bp mRNA linear PRI 29-Apr-1999  
Homo sapiens mirk-related peptide 1 mRNA complete cds.

AF071002.1 GI:4704422

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

TITLE

REFERENCE

AUTHORS

JOURNAL

Human  
Homo sapiens  
Bovine  
Mammalia: Eutheria: Primates: Catarrhini: Hominoidea: Homo.  
1 (bases 1 to 732)  
Timothy K.W., Keating M.T. and Goldstein S.A.  
MIRK forms two potassium channels with HMRG and is associated with  
Cell 92 Activity (2), 175-187 (1999)

2 (bases 1 to 732)  
Abbott G.W., Seethi F., Buck M.E. and Goldstein S.A.N.  
Direct Submission  
Submitted (05-JUN-1998) Section of Developmental Biology and  
Biophysics, Department of Pediatrics and Boyer Center for Molecular

Medicine, Yale University School of Medicine, 295 Congress Avenue,  
New Haven, CT 06536, USA  
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/protein\_id="F4D28086.1"  
/db.xref="GI:4704423"  
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LESSAFTHENIDAKGPKMSF 202 t

BASE COUNT 221 a 152 c 157 g 202 t

ORIGIN

Query Match 100.0%; Score 372; DB 9; Length 732;  
Best Local Similarity 100.0%; Pred. No. 1.3e-100;  
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 ATGCTACTTATTCCTTACCAATTCACAGACGCTGGAGAACGCTCTCCAGAGATTTTAT 60  
|||||  
Db 74 ATGCTACTTATTCCTTACCAATTCACAGACGCTGGAGAACGCTCTCCAGAGATTTTAT 133  
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Oy 61 ACTTATATGACAAATTTGGCCGAGAACACAGACTGAGAGAGAGGCGCTCCAGCCAA 120  
|||||  
Db 134 ACTTATATGACAAATTTGGCCGAGAACACAGACTGAGAGAGAGGCGCTCCAGCCAA 193  
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Oy 121 GTTGATGCTGAGAACTTCTACTATGTCTGCTGACCTGAGATTTGGAATGTC 180  
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Db 194 GTTGATGCTGAGAACTTCTACTATGTCTGCTGACCTGAGATTTGGAATGTC 253  
|||||

Oy 181 TCTTTCATCATGTGGGCGCATCTGCTGAGACCTGTGAATCCAGAGAGGGAACATCC 240  
|||||  
Db 254 TCTTTCATCATGTGGGCGCATCTGCTGAGACCTGTGAATCCAGAGAGGGAACATCC 313  
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Oy 241 AATGACCCCTACACAGATATTTGTGAGAGATGCGCAGGAAAAGTACAGAGCCAAATC 300  
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Oy 301 TTGAACTTGAAGATTCGAAGCCGACACCTGCTGAGAACATTTGTGGGCTGCTTAA 360  
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Oy 361 ATGTGCCCTGTA 372  
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RESULT 2  
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LOCUS Homo sapiens voltage-gated K+ channel subunit MIRP1 (KCNB2) mRNA,  
DEFINITION complete cds.  
ACCESSION AF302095.1 GI:10121897  
VERSION AF302095.1  
KEYWORDS human.  
SOURCE Homo sapiens  
ORGANISM Homo sapiens  
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
REFERENCE 1 (bases 1 to 809)  
TITLE Cloning of human MIRP1 cDNA  
AUTHORS Domenech,A., Estivill,X. and de la Lanza,S.  
JOURNAL Domenech,A., Estivill,X. and de la Lanza,S.  
Direct Submission

JOURNAL  
Submitted (01-SEP-2000) Medical and Molecular Genetics Center,  
Institut Recerca Oncologica, Avda. de Casseldefels Km 2,7,  
L'Hospitalet de Llobregat, Barcelona 08907, Spain  
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BASE COUNT 247 a 172 c 189 g 200 t 1 others

ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.4e-100;  
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Oy 61 ACTTATATGACAAATTTGGCCGAGAACACAGACTGAGAGAGAGGCGCTCCAGCCAA 120  
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Oy 121 GTTGATGCTGAGAACTTCTACTATGTCTGCTGACCTGAGATTTGGAATGTC 180  
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Db 261 GTTGATGCTGAGAACTTCTACTATGTCTGCTGACCTGAGATTTGGAATGTC 320  
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Oy 181 TCTTTCATCATGTGGGCGCATCTGCTGAGACCTGTGAATCCAGAGAGGGAACATCC 240  
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Db 321 TCTTTCATCATGTGGGCGCATCTGCTGAGACCTGTGAATCCAGAGAGGGAACATCC 380  
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Oy 241 AATGACCCCTACACAGATATTTGTGAGAGATGCGCAGGAAAAGTACAGAGCCAAATC 300  
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Db 381 AATGACCCCTACACAGATATTTGTGAGAGATGCGCAGGAAAAGTACAGAGCCAAATC 440  
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Oy 301 TTGAACTTGAAGATTCGAAGCCGACACCTGCTGAGAACATTTGTGGGCTGCTTAA 360  
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Db 441 TTGAACTTGAAGATTCGAAGCCGACACCTGCTGAGAACATTTGTGGGCTGCTTAA 500  
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Oy 361 ATGTGCCCTGTA 372  
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Db 501 ATGTGCCCTGTA 512  
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RESULT 3  
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LOCUS Homo sapiens genomic DNA, chromosome 21q22.1, D21S226-AKL region,  
DEFINITION clone:Q12C8, complete sequence.  
ACCESSION AP000320.1 GI:4835689  
VERSION AP000320.1  
KEYWORDS human.  
SOURCE Homo sapiens  
ORGANISM Homo sapiens  
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
REFERENCE 1 (bases 1 to 24608)  
TITLE Human chromosome 21q22.1  
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
Fujiyama,A., Iwada,K., Tanaka,T., and Sakaki,Y.  
JOURNAL Human chromosome 21q22.1  
Published only in Database (1999) in press

REFERENCE 2 (bases 1 to 24608)  
 AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
 Fujiyama,A., Yada,T., Toki,Y. and Sakaki,Y.  
 JOURNAL Direct Submission (1999) to the DDB/JEM/Genbank database  
 TITLE Mesahira Hattori, The Institute of Physical and Chemical Research  
 (RIKEN), Genomic Sciences Center (GSC), Kitasato Univ., 1-15-1  
 Kitasato, Sagamihara, Kanagawa 228-8555, Japan  
 (E-mail:hattori@ric.riken.go.jp, URL:http://hnp.gsc.riken.go.jp/  
 Tel:81-42-778-9923, Fax:81-42-778-9923)  
 COMMENT The sequence is a part of the data (ACCESSION No. AP000165 -  
 Mesahira Hattori, The Institute of Physical and Chemical  
 Research (RIKEN).  
 The sequencing project is supported by Japan Science Technology  
 Corporation (JST) and The Institute of Physical and Chemical  
 Research (RIKEN).  
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 Oy 121 GTTATGCTGACAACTTCACTATGTATGATCTGATCTGATCTGATCTGATCTGATCTG 180  
 Db 15816 GTTATGCTGACAACTTCACTATGTATGATCTGATCTGATCTGATCTGATCTGATCTG 15875  
 Oy 181 TCTTATCATCTGATGCGCATCTCTGATGACACTGTGAATTCAGAGAGGGACACTCC 240  
 Db 15876 TCTTATCATCTGATGCGCATCTCTGATGACACTGTGAATTCAGAGAGGGACACTCC 15955  
 Oy 241 AATGACCCCTACACACAGTATCTGTGAGAGACGTGGACGAAAGATGACACCAATTC 300  
 Db 15936 AATGACCCCTACACACAGTATCTGTGAGAGACGTGGACGAAAGATGACACCAATTC 15995  
 Oy 301 TTGAATCTAGAGAAATCGAAGGCCCATCATCATGAGAAACATTTGTCGCGTGGTCAA 360  
 Db 15996 TTGAATCTAGAGAAATCGAAGGCCCATCATCATGAGAAACATTTGTCGCGTGGTCAA 16055  
 Oy 361 ATGTCCCTCTGA 372  
 Db 16056 ATGTCCCTCTGA 16067  
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 LOCUS AP000052 100000 bp DNA linear PRI 20-NOV-1999  
 DEFINITION Homo sapiens genomic DNA, chromosome 21q22.1, segment 23/28,  
 complete sequence.  
 ACCESSION AP000052  
 VERSION AP000052.1 GI:1132362  
 SOURCE Homo sapiens DNA, clone:245P17-f4A4f\_2.  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 REFERENCE 1 (bases 1 to 100000)  
 AUTHORS Hattori,M., Ishii,K., Toyoda,A., Shiba,T. and Sakaki,Y.  
 TITLE Homo sapiens genomic DNA, chromosome 21q

JOURNAL Published Only in Database (1998) in press  
 REFERENCE 2 (bases 1 to 100000)  
 AUTHORS Hattori,M., Ishii,K., Toyoda,A., Shiba,T. and Sakaki,Y.  
 JOURNAL Direct Submission (1998) to the DDB/JEM/Genbank database  
 TITLE Mesahira Hattori, Kitasato University of Science, JST  
 Sequencing Laboratory, Kitasato 1-15-1, Sagamihara 228, Japan  
 (E-mail:hattori@ric.riken.go.jp, Tel:0427-78-9732,  
 Fax:0427-78-9561)  
 COMMENT This sequence is conducted by Kitasato University JST sequencing  
 Laboratory as a JST sequencing team.  
 Principal Investigator: Yoshinuki Sakaki Ph.D.  
 sakaki@ipc.ims.u-tokyo.ac.jp \*81-3-5449-5445.  
 Sub-leader: Tadayoshi Shiba Ph.D., Mesahira Hattori Ph.D. The  
 sequence is submitted by Human Genome Sequencing in AUs project of  
 JST  
 Japan Science and Technology Corporation (JST)  
 5-3, Yonbancho, Chiyoda-ku, Tokyo 102-0028 Japan  
 For further information about this sequence, including its location  
 on chromosome 21, please visit the following web site:  
 archive Web site (http://www.ahis.tokyo.jst.go.jp/BS/top.html)  
 or send email to webmaster@www.ahis.tokyo.jst.go.jp/;.  
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 Oy 121 GTTATGCTGACAACTTCACTATGTATGATCTGATCTGATCTGATCTGATCTGATCTG 180  
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 Oy 181 TCTTATCATCTGATGCGCATCTCTGATGACACTGTGAATTCAGAGAGGGACACTCC 240  
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 Oy 241 AATGACCCCTACACACAGTATCTGTGAGAGACGTGGACGAAAGATGACACCAATTC 300  
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 Oy 361 ATGTCCCTCTGA 372  
 Db 80578 ATGTCCCTCTGA 80589  
 RESULT 5  
 LOCUS AP000120 100000 bp DNA linear PRI 25-SEP-1999  
 DEFINITION Homo sapiens genomic DNA of 21q22.1, GAT and AAT related,  
 SL53A3-f4A4 region, segment 3/8, complete sequence.  
 ACCESSION AP000120  
 VERSION AP000120.1 GI:4730889

KEYWORDS HTG.  
SOURCE Homo sapiens DNA.  
ORGANISM Homo sapiens.  
REFERENCE Hattori, M., Ishii, K., Toyoda, A., Shiba, T., and Sakaki, Y.  
AUTHORS Hattori, M., Ishii, K., Toyoda, A., Shiba, T., and Sakaki, Y.  
TITLE Homo sapiens 817,199bp genomic DNA of 21q22.1 GANT and AML region  
JOURNAL Published Only in Databases (1999) in press  
REFERENCE 2 (bases 1 to 100000)  
AUTHORS Hattori, M., Ishii, K., Toyoda, A., Shiba, T., and Sakaki, Y.  
TITLE Direct Submission  
JOURNAL Submitted (15-APR-1999) to the DDBJ/EMBL/Genbank databases, Mita  
Hitachi, Japan Science and Technology Corporation (JST), Advanced  
Databases Department: 5-3, Yonbancho, Chiyoda-ku, Tokyo 102-0081,  
Japan (E-mail: mtk@res.tokyo.jst.go.jp/  
URL: http://www.a11s.tokyo.jst.go.jp/, Tel: 81-3-5214-8491,  
Fax: 81-3-5214-8470)  
COMMENT The JST is conducted by Kitano University JST sequencing  
laboratory as a JST sequencing team  
Principal Investigator: Yoshiaki Sakaki Ph.D.  
Phone: +81-3-5449-5622, Fax: +81-3-5449-5445,  
sakaki@jst.tokyo.ac.jp  
Sub-leader: Tadayoshi Shiba Ph.D., Masahiro Hattori Ph.D. The  
sequence is submitted by Human Genome Sequencing in AUs project of  
JST, Science and Technology Corporation (JST)  
5-3 Yonbancho, Chiyoda-ku, Tokyo 102-0081 Japan  
For further information about this sequence, including its location  
and relationship to other sequences, please visit our sequence  
archive Web site (http://www.a11s.tokyo.jst.go.jp/HGS/) or send  
email to webmaster@www.a11s.tokyo.jst.go.jp.  
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Best Local Similarity 100.0%; Pval: No. 3.9e-100;  
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Db 89760 AATGACCTTACAGACAGATTTGAGAGAGCTGGCGAGCAAAAAGTACAGACCCAAATC 89819  
Oy 301 TTGATATGAGCAATTTGGCGGAGAACACAGACCTGCGACAGCCCTCAAGCCAA 360  
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Db 89880 ATGTCCTCCCTGA 89891  
RESULT 5  
AP000167 100000 bp DNA linear. PRI: 08-JAN-2000  
LOCUS Homo sapiens genomic DNA, chromosome 21q22.1, D21S226-AML region,  
clone B2344741-F5088, Hsegant 3/9, complete sequence.  
DEFINITION  
ACCESSION AP000167.1 GI:4827132  
VERSION  
KEYWORDS HTG.  
SOURCE Homo sapiens DNA.  
ORGANISM Homo sapiens.  
REFERENCE Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,  
AUTHORS Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,  
Fujiyama, A., Yada, T., Tokoki, Y., and Sakaki, Y.  
TITLE Homo sapiens 8234471-F5088  
JOURNAL Published Only in Databases (1999) in press  
REFERENCE 2 (bases 1 to 100000)  
AUTHORS Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,  
Fujiyama, A., Yada, T., Tokoki, Y., and Sakaki, Y.  
TITLE Direct Submission (1999) to the DDBJ/EMBL/Genbank databases  
Masahiro Hattori, The Institute of Physical and Chemical Research  
(RIKEN), Genomic Science Center (GSC), Kitasato Univ., 1-15-1  
Kitasato, Sagamihara, Kanagawa 228-8555, Japan  
(E-mail: hattori@psc.riken.go.jp, URL: http://hnp.gsc.riken.go.jp/  
Tel: 81-42-778-9923, Fax: 81-42-778-9924)  
COMMENT The present data does not contain E.  
coli transposon insertion. The present data does not contain E.  
coli transposon insertion. The present data does not contain E.  
original/previous sequences. We determined the boundary between  
the insertion and genomic sequences experimentally. The sequencing  
project is supported by Japan Science Technology Corporation (JST)  
and The Institute of Physical and Chemical Research (RIKEN).  
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Oy 181 TCTTATATGAGCAATTTGGCGGAGAACACAGACCTGCGACAGCCCTCAAGCCAA 240  
Db 72365 TCTTATATGAGCAATTTGGCGGAGAACACAGACCTGCGACAGCCCTCAAGCCAA 240  
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Oy 121 GTTATGTCGACAACTCTACATCTGATACACCTGACACCTGACACCTGACACCTGACAAATGTC 180
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Db 301547 GTTATGTCGACAACTCTACATCTGATACACCTGACACCTGACACCTGACACCTGACAAATGTC 301606
Oy 181 TCTTTCATCATCTGCGGCGCATCTGCTGTACACATGTGAAATCCAAAGACGCAACATCC 240
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Db 301607 TCTTTCATCATCTGCGGCGCATCTGCTGTACACATGTGAAATCCAAAGACGCAACATCC 301666
Oy 241 AATGACCCCTACGACAGCAATATTTGAGAGAGCTGGCGAGGAAAGTACAAAGACCAATC 300
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Db 301667 AATGACCCCTACGACAGCAATATTTGAGAGAGCTGGCGAGGAAAGTACAAAGACCAATC 301726
Oy 301 TTGAATCTAGAGAAATGCAAGGCGCAACATCCATGTGAAACATTTGCTGCTGGGTTCAAA 360
Db 301727 TTGAATCTAGAGAAATGCAAGGCGCAACATCCATGTGAAACATTTGCTGCTGGGTTCAAA 301786
Oy 361 ATGTCCCTCTGA 372
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Db 301787 ATGTCCCTCTGA 301798

BESTIT 8
LOCUS BC022699 1664 bp mRNA linear ROD 04-FEB-2002
DEFINITION Mus musculus, RIKEN cDNA 2200002j16 gene, clone MGC:31447
IMAGE:4481325, mMAN, complete cds.
ACCESSION BC022699
VERSION BC022699.1 GI:18490550
KEYWORDS MGC, EST, mRNA, complete cds.
SOURCE Mus musculus
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Scuriognathi; Muridae; Murinae; Mus.
1 (bases 1 to 1664)
REFERENCE
Strasberg/R.
Direct Submission
Submitted (01-FEB-2002) National Institutes of Health, Mammalian
Genome Project, Bethesda, MD
Accession: BC022699.1
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2850,
USA
NIH-MGC Project URL: http://mgc.ncl.nih.gov
Contact: MGC help desk
Email: gcraps-remail.nih.gov
Tissue Procurement: The Cyto Laboratory
Genome Project, 1000 University Ave., 10th floor,
CDNA Library created by the I.M.A.G.E. Consortium (ILM)
DNA sequencing by: Sequencing Group at the Stanford Human Genome
Center, Stanford University School of Medicine, Stanford, CA 94305
Web site: http://www-sngc.stanford.edu
Contact: (Dickson, Mark) mcdickpax1.stanford.edu
Dickson, M., Schmutz, J., Grimwood, J., Rodriguez, A., and Myers,
R. M.
Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/ILM at: http://image.llnl.gov
Series: IRAK plate: 44 Row: a Column: 22
This clone was selected for full length sequencing because it
passed the following selection criteria: Hexamer frequency ORF
analysis.
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Oy	269	TAGCATCTGGAGGAAAATACTGAC	291	
Db	388	AACGACATCTGGACAGCAAAGACAAGC	411	
RESULT 14				
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DEFINITION	Cavia porcellus potassium channel mRNA, complete cds.		R0D 11-MAY-1994	
ACCESSION	L20462.1			
VERSION	L20462.1 GI:484140			
KEYWORDS	potassium channel;			
SOURCE	Cavia porcellus adult cardiac muscle cDNA to mRNA.			
ORGANISM	Cavia porcellus			
REFERENCE	Bukharjaya; Metacoss; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Artiodactyla; Rodentia; Hystriognathii; Cavidae; Cavia. 1 (bases 1 to 750)			
AUTHORS	Zhang,J., Jurkiewicz,N.K., Polander,K., Lazarides,E., Salate,J.J. and Swanson,R.			
TITLE	K+ currents expressed from the guinea pig cardiac IsK protein are enhanced by activators of protein kinase C			
JOURNAL	Proc. Natl. Acad. Sci. U.S.A. 91, 1766-1770 (1994)			
FEATURES	Location/Organisms			
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BASE COUNT	189 a 193 c 114 g 108 t			
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Best Local Similarity	66.4%; Pred. No. 1.4e-05;			
Matches	95; Conservative 0; Mismatches 45; Indels 3; Gaps 1;			
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Oy	211 ACTGGGAATCCCAGAAGCAAGGACAACTCTCAATGACCCCTCACACACAGTAGTAACTTTGTAG--	TA 267		
Db	255 TATATTCGATCCAGAAACCTGGGAGCACTGCGACGACCGTCCTCAAGGTGATCATCGAGTCA	314		
Oy	268 GAGCATGCGAGAGAAAAATGACA	290		
Db	315 GACACTGCGAAGAAAGACAA	337		
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GPIISK		390 bp	mRNA	linear
DEFINITION	Homo sapiens delayed rectifier potassium channel subunit ISK mRNA, complete cds.		PRI 14-APR-1995	
ACCESSION	AF135188			
VERSION	AF135188.1 GI:4583498			
KEYWORDS	.			
SOURCE	human.			
ORGANISM	Homo sapiens			

	REFERENCE	Eukaryotes; Metazoa; Chordata; Craniata; Vertebrates; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OY	AUTHORS	Rae,J.L., rectifier potassium channel subunit from human cornea
Dc	JOURNAL TITLE	biochim biophys acta
OY	JOURNAL	unpublished
Dc	RECEIVED	2 (bases 1 to 390)
OY	REFERENCE TITLE	Rae,J.L.
Dc	JOURNAL	Submitted (16-MAR-1999) Department of Physiology and Biophysics, Mayo Foundation, 200 1st Street SW, Rochester, MN 55905, USA
	FEATURES	Location/Qualifiers
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	primer_bind	complement(372..390)
	BASE COUNT	123 c 102 g 75 t
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OY	210	CATTGTGAATCAAGAACAAGGCAATCTCAATGATGATGCCATCCATCCATCCATGATGATC-- 266
Dc	192	CTCATCTCCCTCCATGAAGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 251
OY	267	AGAGACATGCGAGAAAAGATACAGAGCCATATC 300
Dc	252	CGATGCTGCGCACAGAACAGATAGAGCGCTATGTC 285

Search completed: October 24, 2002, 13:54:33  
Job time : 2063 secs

Job time : 2063 secs

GenCore version 5.1.3  
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 24, 2002, 13:14:11 Search time 198 Seconds  
(without alignments)

3225.718 Million cell updates/sec

Title: US-09-550-163-1-COPY\_74\_445

Sequence: 1 agctcacttaccattc.....ggtccaaatgccccccgga 372

Scoring table: IDENTITY-MDC

Gapop 10.0 , Capext 1.0

Searched: 1736436 seqs, 858457221 residues

Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 43 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	372	100.0	372	22	AA124432	Probe #14365 for g
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4	372	100.0	471	22	AA809192	Nucleic acid sequen
5	372	100.0	600	22	AA809192	Human M18P1 domo
6	372	100.0	600	22	AA852645	Human polyunoclel
7	372	100.0	655	22	AA851661	Human polyunoclel
8	372	100.0	732	21	AA64071	Human potassium ch
9	370.4	99.6	732	21	AA64083	Human potassium ch

10	370.4	99.6	732	21	AA64084	Human potassium ch
11	370.4	99.6	732	21	AA64085	Human potassium ch
12	370.4	99.6	732	21	AA64086	Human potassium ch
13	312	83.9	312	22	AA84938	Human breast cell
14	312	83.9	312	22	AA867836	Human foetal liver
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17	312	83.9	312	22	AA867836	Human foetal liver
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33	53.2	14.3	231	22	AA868478	Human foetal liver
34	53.2	14.3	231	22	AA868478	Human foetal liver
35	53.2	14.3	231	22	AA868478	Human foetal liver
36	53.2	14.3	231	22	AA868478	Human foetal liver
37	53.2	14.3	231	22	AA868478	Human foetal liver
38	53.2	14.3	231	22	AA868478	Human foetal liver
39	53.2	14.3	231	22	AA868478	Human foetal liver
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#### ALIGNMENTS

RESULT 1  
AA124432  
ID: AA124432 standard; DNA, 372 BP.

NC  
AA124432:  
12-CT-2001 (first entry)

DT  
Probe #14365 for gene expression analysis in human cervical cell sample.

DE  
Probe: human; microarray; gene expression; cervical epithelial cell;

DE  
cervical cancer; ss.

KM  
Homo sapiens.

OS  
Homo sapiens.

PN  
MO20157278-A2.

XX  
09-RUC-2001.

XX  
30-JAN-2001; 2001MO-0500670.

XX  
04-FEB-2000; 2000US-0180312.

XX  
26-MAY-2000; 2000US-0207456.

XX  
30-JUN-2000; 2000US-0608408.

XX  
03-RUC-2000; 2000US-0652265.

XX  
27-SEP-2000; 2000US-0235350.

XX  
04-CT-2000; 2000CB-0024363.

XX  
(NOTE-) MOLECULAR DYNAMICS INC.

XX  
Penn SG, Hanzel DK, Chen W, Rank DR;

XX WP1: 2001-468901/53.  
 XX Human genome-derived single exon nucleic acid probes useful for  
 XX analyzing gene expression in human cervical epithelial cells -  
 XX  
 XX Claim 25: SEQ ID No 1365; 48Pp; English.  
 XX  
 XX The present invention relates to human single exon nucleic acid probes  
 XX for the present sequence. One such probe, the 58Ns are derived  
 XX from human cervical cells and are useful for measuring human gene  
 XX expression, which can be used for measuring human gene expression in a  
 XX sample derived from human cervical epithelial cells. By measuring gene  
 XX expression, the probes are therefore useful in grading and/or staging  
 XX of diseases of the cervix, notably cervical cancer.  
 XX Note: The sequence data for this patent did not form part of the printed  
 XX publication of the patent document in the format directly from WIPO  
 XX at ftp://wipo.int/pub/published\_pat\_sequences.  
 XX  
 XX Sequence 372 Bp: 110 A; 90 C; 82 G; 90 T; 0 other;  
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 XX Best Local Similarity 100.0%; Pred. No. 1.6e-102;  
 XX Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
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 XX RESULT 2  
 XX ID AA109965  
 XX AA109965 standard; DNA: 372 Bp.  
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 XX AA109965:  
 XX  
 XX 09-OCT-2001 (first entry)  
 XX  
 XX Probe #9956 used to measure gene expression in human breast sample.  
 XX  
 XX Probs: human; breast disease; breast cancer; development disorder; ss;  
 XX Inflammatory disease; proliferative breast disease; non-carcinoma tumour.  
 XX  
 XX Homo sapiens.  
 XX  
 XX MO200157270-A2.  
 XX  
 XX 09-AUG-2001.  
 XX  
 XX

XX 29-JAN-2001; 2001WO-US00661.  
 XX  
 XX 04-FEB-2000; 2000US-1493132.  
 XX PR 29-JAN-2000; 2000US-0207456.  
 XX PR 30-JUN-2000; 2000US-0608408.  
 XX PR 03-AUG-2000; 2000US-0632366.  
 XX PR 21-SEP-2000; 2000US-0234687.  
 XX PR 27-SEP-2000; 2000US-0236359.  
 XX PR 04-OCT-2000; 2000US-0024263.  
 XX  
 XX (MOLE-) MOLECULAR DYNAMICS INC.  
 XX  
 XX Penn Sq, Hanzel Dk, Chen W, Rank Dr.  
 XX  
 XX WP1: 2001-476286/51.  
 XX  
 XX Novel single exon nucleic acid probe used to measuring gene expression  
 XX in a human breast -  
 XX  
 XX Claim 25: SEQ ID No 9956; 322pp; English.  
 XX  
 XX The present invention relates to novel single exon nucleic acid probes.  
 XX The present sequence is one such probe. The probes are useful for  
 XX measuring human gene expression in a human breast sample, where the probe  
 XX hybridises at high stringency to a nucleic acid expressed in the human  
 XX breast, and is useful for measuring human gene expression in the human  
 XX breast, monitoring and prognosing diseases of the human breast,  
 XX particularly those diseases with polygenic aetiology. The diseases  
 XX include: breast cancer, disorders of development, inflammatory diseases  
 XX of the breast, fibrocystic changes, proliferative breast disease and  
 XX non-carcinoma tumours.  
 XX Note: The sequence data for this patent did not form part of the printed  
 XX publication of the patent document in the format directly from WIPO  
 XX at ftp://wipo.int/pub/published\_pat\_sequences.  
 XX  
 XX Sequence 372 Bp: 110 A; 90 C; 82 G; 90 T; 0 other;  
 XX  
 XX Query Match 100.0%; Score 372; DB 22; Length 372;  
 XX Best Local Similarity 100.0%; Pred. No. 1.6e-102;  
 XX Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
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 XX 361 ATGTCCCCCTGGA 372  
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 XX RESULT 3  
 XX ID AAS00245  
 XX AAS00245 standard; DNA: 372 Bp.  
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XX  AAS00245:
AC  10-MAY-2001 (first entry)
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DX  Human potassium channel regulatory protein, Mink2, DNA sequence.
XX
XX  Human; Mink2; potassium channel; cardiac arrhythmia; hypertension; ds;
KM  angina; asbestia; diabetes; renal insufficiency; urinary incontinence;
XX  irritable colon; epilepsy; cerebrovascular ischemia; autoimmune disease.
XX
XX  Homo sapiens.
XX
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XX  FH      1..372
XX  CDS      /'tag' = a
XX           /product= "MINK2 potassium channel protein"
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XX  W0200114403-A1.
XX
XX  01-MAR-2001.
XX
XX  18-AUG-2000; 2000MO-0522799.
XX
XX  20-AUG-1999; 99US-0379201.
XX
XX  (UYCA-) UNIT CASE WESTERN RESERVE.
XX
XX  Flicker E., Mable B., Brown AM.
XX
XX  WPI: 2001-218424/22.
XX  P-PSDB: AA000215.
XX
XX  Novel potassium channel gene termed Mink2 encoding potassium channel
XX  regulatory protein, useful for screening compounds that are useful for
XX  treating diseases caused by aberrant potassium activity -
XX
XX  Claim 1; Fig 9; 39pp; English.
XX
XX  The sequence represents the coding sequence of human potassium channel
XX  regulatory protein, Mink2. Mink2 sequence is useful for producing a
XX  potassium channel regulatory protein useful for in vitro or in vivo
XX  screening of agonistic or antagonistic compounds that are useful for
XX  treating diseases caused by aberrant potassium activity, such as human
XX  cardiac arrhythmias, hypertension, urinary incontinence, irritable
XX  colon, epilepsy, cerebrovascular ischemia, and autoimmune disease.
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XX  Best Local Similarity 100.0%; P-val: 1e-102;
XX  Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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XX  DB  61 ACTTATATGAGCATTTGGCGCCAGAACACAGACGTGAGCAAGGCGCTTCACAGCAA 120
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XX  ID AAF80269 standard; DNA; 471 BP.
XX  XX
XX  AAF80269;
XX  DT 29-JUN-2001 (first entry)
XX
XX  Nucleotide sequence of human potassium channel subunit ISK2.
XX
XX  Human; potassium channel; ISK2; gene therapy; gastric motility;
XX  gastric acid secretion; anti-arrhythmic agent; myocardial infarction; ss.
XX
XX  Homo sapiens.
XX
XX  Key      Location/Qualifiers
XX  FH      79..450
XX  CDS      /'tag' = a
XX           /product= "potassium channel subunit ISK2"
XX
XX  W0200127246-A1.
XX
XX  19-APR-2001.
XX
XX  10-OCT-2000; 2000MO-0528014.
XX
XX  12-OCT-1999; 99US-0158781.
XX
XX  (MERI) MERCK & CO INC.
XX
XX  Swanson RJ, Liu Y, Polander K.
XX
XX  WPI: 2001-273764/28.
XX  P-PSDB: AAB67800.
XX
XX  New DNA encoding the ISK2 potassium channel subunit, useful e.g. for
XX  detecting mutations and screening for therapeutic agents -
XX
XX  Claim 3; Fig 1A; 46pp; English.
XX
XX  The present sequence encodes a human potassium channel subunit,
XX  designated ISK2, the cDNA polymethionide, and derived probes,
XX  useful for identifying and detecting the presence of ISK2 gene
XX  levels of mRNA expression and to isolate homologous sequences; for
XX  recombinant expression of ISK2; in gene therapy to increase potassium
XX  channel activity and to generate transgenic animals, as models and
XX  for drug screening. Recombinant ISK2 is used for studying biochemical
XX  activity of ISK2 and its role in disorders of gastric motility and
XX  gastric acid secretion, and to raise specific antibodies. ISK2
XX  is used for screening for compounds that increase or decrease
XX  increased or reduced potassium channel activity e.g. as
XX  anti-arrhythmic agents for treating myocardial infarction and as
XX  regulators of gastric acid secretion.
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XX  Sequence 471 BP; 143 A; 110 C; 103 G; 115 T; 0 other:
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XX  Best Local Similarity 100.0%; P-val: 1e-102;
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XX      |||
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AC	AAK52645;
ID	AAK52645 standard; cDNA: 600 bp.
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XX	06-NOV-2001 (first entry)
DE	human polynucleotide seq ID NO 2174.
KW	human; cytokine; cell proliferation; cell differentiation; gene therapy;
KW	vaccine; peptide therapy; stem cell growth factor; hematopoiesis;
KW	tissue growth factor; immunomodulator; cancer; leukemia;
XX	neuro system disorder; arthritis; inflammation; ss.
XX	
XX	Homn sapiens.
XX	
XX	WO200157150-A2.
XX	
XX	09-AUG-2001.
XX	
XX	05-FEB-2001; 2001MO-US04098.
XX	
XX	03-FEB-2000; 2000US-0496914.
XX	27-APR-2000; 2000US-0560875.
XX	20-JUN-2000; 2000US-0598075.
XX	19-JUL-2000; 2000US-0620325.
XX	01-SEP-2000; 2000US-0654936.
XX	15-SEP-2000; 2000US-0653661.
XX	20-OCT-2000; 2000US-0693325.
XX	30-NOV-2000; 2000US-0728422.
XX	
XX	(HYSE)- HXSD INC.
XX	
XX	Tang YT, Liu C, Dmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;
XX	Zhao QA, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZM;
XX	Xue AJ, Yang Y, Wejhrman T, Goodrich R;
XX	
XX	WPI: 2001-476283/51.
XX	F-PDB; AAK79312.
XX	
XX	Nucleic acids encoding polypeptides with cytokine-like activities,
XX	useful in diagnosis and gene therapy -
XX	
XX	Claim 1: Page 4539-4540: 6221pp: English.
XX	
XX	The invention relates to polynucleotides (AAK51456-AAK53435) and the
XX	encoded polypeptides (AAK79312-AAK80302) that exhibit activity elating to
XX	proliferation, cell proliferation or cell differentiation or which may induce
XX	proliferation, cell proliferation or cell differentiation. The vaccines or
XX	polynucleotides and polypeptides are useful in gene therapy or
XX	peptide therapy. The polypeptides have various cytokine-like activities,
XX	e.g. stem cell growth factor activity, hematopoiesis regulating
XX	activity, tissue growth factor activity, immunomodulator activity and
XX	treatment of cancer, leukemia, nervous system disorders, arthritis and
XX	inflammation.
XX	For SEQ ID NO 2110 (AAK52861), 2111 (AAK52882) and 3666
XX	(AAK80020) the omitted nucleotide pages from the sequence listing
XX	were missing at the time of publication.
XX	
XX	Sequence 600 bp: 187 A; 133 C; 130 G; 144 T; 6 other:
XX	
XX	Query Match 100.0%; Score 172; DB 22; Length 600;
XX	Best Local Similarity 100.0%; Pred. No. 2e-102;
XX	Matches 572; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX	
0Y	1 AGCTAGCTATTCGCAATTTTCAGACAGAGCGTCGACGAGATTTTAT 60
0Y	38 ATGCTAGCTATTCGCAATTTTCAGACAGAGCGTCGACGAGATTTTAT 97
0Y	61 ACTTATATGCAATTTGCGCGCAGAAACACGAGTCGACGAGGCGTCGACGACAA 120

Db	98	ACTTAAATGACAAATTGGGCGCGAGAAACAAACAGCTGAGAGGAGGCGCTCCCAACCAACAA	157
Qy	121	GTGATGCTGACAAACTGCTACCTACCTATGTCATCTGCTGACTCATGCTGATGATGTAATGTC	180
Db	158	GTGATGCTGACAAACTGCTACCTACCTATGTCATCTGCTGACTCATGCTGATGATGTAATGTC	217
Qy	181	TCCTTATCATCTGCGGCGCATCTCGGTGAGCACTGCTGAATTCAGAGAGAGGAGCAACCTCC	240
Db	218	TCCTTATCATCTGCGGCGCATCTCGGTGAGCACTGCTGAATTCAGAGAGAGGAGCAACCTCC	277
Qy	241	AATGACCCCTGACACCCAGTACATGTTGATGAGAGCTGCGACGAGAAATGACAGACCAATTC	300
Db	278	AATGACCCCTGACACCCAGTACATGTTGATGAGAGCTGCGACGAGAAATGACAGACCAATTC	337
Qy	301	TTCGATCTGACAGAAATGACAGGCGCACATCCATGCTATGACAAATCTGGCGGCTGCGTTC	350
Db	338	TTCGATCTGACAGAAATGACAGGCGCACATCCATGCTATGACAAATCTGGCGGCTGCGTTC	397
Qy	361	ATGTCCCTGCA 372	
Db	398	ATGTCCCTGCA 409	
RESULT 7			
AAKS1661			
ID	AAKS1661	standard; cDNA; 655 BP.	
XX	AAKS1661.		
DT	06-NOV-2001	(first entry)	
DE	Human polynucleotide SEQ ID NO 206.		
KM	Human; cytokine; cell proliferation; cell differentiation; gene therapy;		
KM	vaccine; peptide therapy; stem cell growth factor; haemotopoiesis;		
KM	neurotic system disorder; arthritis; inflammation; sbs.		
XX	Homo sapiens.		
XX	MO200157190-A2.		
PN	09-AUG-2001.		
PR	05-FEB-2001; 2001MO-US04098.		
PR	03-FEB-2000; 2000US-0496934.		
PR	27-APR-2000; 2000US-0560875.		
PR	20-JUN-2000; 2000US-0596075.		
PR	19-JUL-2000; 2000US-0620325.		
PR	01-SEP-2000; 2000US-0654936.		
PR	13-SEP-2000; 2000US-0654936.		
PR	20-OCT-2000; 2000US-0693125.		
PR	30-NOV-2000; 2000US-0728422.		
PA	(HYSE-) HYSEQ INC.		
PI	Teng YF, Liu C, Dumanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;		
PI	Zhao QH, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;		
PI	Xue AJ, Teng Y, Wejntman T, Goodrich R;		
PI	WPI: 2001-476283/51.		
DR	P-PSDB: AAT78528.		
PT	Nucleic acids encoding polypeptides with cytokine-like activities,		
PT	useful in diagnosis and gene therapy -		
XX	Claim 1; Page 1024; 6221pp; English.		
CC	The invention relates to polynucleotides (AAKS1456-AAKS5435) and the		
CC	encoded polypeptides (AAW8323-AAW8302) that exhibit activity elating to		
CC	cytokine, cell proliferation or cell differentiation or which may induce		

production of other cytokines in other cell populations. The polyclonal antibodies used in these studies are the vaccines or peptide therapy. The polypeptides have various cytotoxic activities, e.g. stem cell growth factor activity, immunomodulatory activity and activity/inhibin activity and may be useful in the diagnosis and/or treatment of cancer, leukemia, nervous system disorders, arthritis and CC Notes: Search for SPO ID NO 2110 (AA55581), 2111 (AA55582) and 3666 (AA66020) are omitted as the relevant pages from the sequence listing were missing at the time of publication.

Sequence 655 BP; 196 A; 154 G; 146 G; 153 T; 6 other;

Query Match 100.0%; Score 372; DB 22; Length 655;

Best Local Similarity 100.0%; P-Id: 2e-10; Mismatches 0; Indels 0; Gaps 0;

Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

0y 1 ATGCTCATTTATTCATTCACACAGCCGGAGAGAGCTCTCCAGAGATTTTATT 60  
Db 93 ATGCTCATTTATTCATTCACACAGCCGGAGAGAGCTCTCCAGAGATTTTATT 152  
0y 61 ACTATATGACATTTGGCCGACAGACACACTGAGAGAGAGCCCTCCAGACAA 120  
Db 153 ACTATATGACATTTGGCCGACAGACACACTGAGAGAGAGCCCTCCAGACAA 212  
0y 121 GTTATGATGAGAACTTACTATGCTACCTGACCTGAGAGAGATTTGATTTTC 180  
Db 213 GTTATGATGAGAACTTACTATGCTACCTGACCTGAGAGAGATTTGATTTTC 272  
0y 181 TTTTATGATGAGAACTTACTATGCTACCTGACCTGAGAGAGATTTGATTTTC 240  
Db 273 TTTTATGATGAGAACTTACTATGCTACCTGAGAGAGATTTGATTTTC 332  
0y 241 AATGACCCCTACACACAGTATGAGAGAGCTGCGAGAGAAAGTCAAGACCAATC 300  
Db 333 AATGACCCCTACACACAGTATGAGAGAGCTGCGAGAGAAAGTCAAGACCAATC 392  
0y 301 TTGAATCTGAGAAATGAGAGAGAGCTGCGAGAGAAAGTCAAGACCAATC 360  
Db 393 TTGAATCTGAGAAATGAGAGAGAGCTGCGAGAGAAAGTCAAGACCAATC 452  
0y 361 ATGTCCCCCTGA 372  
Db 453 ATGTCCCCCTGA 464

RESULT 8  
AAC64071  
ID AAC64071 standard; cDNA; 732 BP.

XX AAC64071;  
XX  
XX  
XX 19-FEB-2001 (first entry)  
XX  
XX Human potassium channel protein KCNE2 (MIRP2) cDNA, SPO ID NO.1.  
XX  
XX Human: KCNE2; MIRP2; potassium channel protein; KCNE2-related;  
XX MIRK-related; long QT syndrome; cardiac arrhythmia;  
XX drug screening; knockout mouse; transgenic animal; ion channel disorder;  
XX HIRG; #6.  
XX  
XX Homo sapiens.  
XX  
XX W020006344-A1.  
XX  
XX 26-OCT-2000.  
XX  
XX 14-APR-2000; 2000MO-US10004.  
XX  
XX 15-APR-1999; 990US-0129404.  
XX

PA (UTAR ) UNIT UTAR RES FOUND.

PA (UTAR ) UNIT TALE.

XX Abbott CW, Seftl F, Splawski I, Keating KR, Goldstein SM:

DR WPI: 2000-672747/65.

XX P-PSDB; AA529585.

XX Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for  
XX drug screening and treating ion channel disorders, especially long QT  
XX syndrome.

XX Claim 1; Page 118-119; 132pp; English.

XX The invention relates to novel ion channel proteins related to  
XX KCNE2 (MIRP2) and to nucleic acids encoding them. The proteins of  
XX KCNE2 (MIRP2) are human MIRP2, MIRP1, MIRP3, MIRP4 and MIRP5,6,  
XX respectively. The human MIRP2 (MIRP2) is encoded by the MIRP2  
XX gene, and human MIRP4 (MIRP4) is encoded by the MIRP4 gene,  
XX respectively. The cDNAs encoding these proteins are given in AAC64071-  
XX AAC64076. KCNE2, along with HIRG, forms cardiac fast delayed rectifier  
XX potassium channels (I-KR), mutations in which are associated with long  
XX QT syndrome. The invention also relates to methods of diagnosing long QT  
XX syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
XX disruption of an endogenous KCNE2 or KCNE3 gene, a transgenic  
XX animal expressing a transgenic animal comprising human KCNE2 and HIRG  
XX cDNA, and methods of and screening drugs for treating long QT syndrome  
XX using KCNE2 proteins (including mutants), nucleic acids encoding them  
XX and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
XX acids, and proteins may be used for diagnosing or treating ion channel  
XX disorders, especially long QT syndrome, transgenic animals comprising  
XX long QT syndrome, and screening drugs for treating long QT syndrome.  
XX The present sequence represents cDNA encoding human KCNE2 (MIRP2).

Sequence 732 BP; 221 A; 152 G; 157 G; 202 T; 0 other;

Query Match 100.0%; Score 372; DB 21; Length 732;

Best Local Similarity 100.0%; P-Id: 2e-10; Mismatches 0; Indels 0; Gaps 0;

Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

0y 1 ATGCTCATTTATTCATTCACACAGCCGGAGAGAGCTCTCCAGAGATTTTATT 60  
Db 74 ATGCTCATTTATTCATTCACACAGCCGGAGAGAGCTCTCCAGAGATTTTATT 133  
0y 61 ACTATATGACATTTGGCCGACAGACACACTGAGAGAGAGCCCTCCAGACAA 120  
Db 134 ACTATATGACATTTGGCCGACAGACACACTGAGAGAGAGCCCTCCAGACAA 193  
0y 121 GTTATGATGAGAACTTACTATGCTACCTGACCTGAGAGAGATTTGATTTTC 180  
Db 194 GTTATGATGAGAACTTACTATGCTACCTGACCTGAGAGAGATTTGATTTTC 253  
0y 181 TTTTATGATGAGAACTTACTATGCTACCTGACCTGAGAGAGATTTGATTTTC 240  
Db 254 TTTTATGATGAGAACTTACTATGCTACCTGAGAGAGATTTGATTTTC 313  
0y 241 AATGACCCCTACACACAGTATGAGAGAGCTGCGAGAGAAAGTCAAGACCAATC 300  
Db 314 AATGACCCCTACACACAGTATGAGAGAGCTGCGAGAGAAAGTCAAGACCAATC 373  
0y 301 TTGAATCTGAGAAATGAGAGAGAGCTGCGAGAGAAAGTCAAGACCAATC 360  
Db 374 TTGAATCTGAGAAATGAGAGAGAGCTGCGAGAGAAAGTCAAGACCAATC 433  
0y 361 ATGTCCCCCTGA 372  
Db 434 ATGTCCCCCTGA 445

RESULT 9  
AAC64083  
ID AAC64083 standard; DNA; 732 BP.

XX AAC64083;  
AC  
XX 19-FEB-2001 (first entry)  
XX  
DE Human potassium channel protein KCNE2 (MIRP1) Q98 mutant DNA.  
XX  
KW Human: KCNE2; MIRP1; potassium channel protein; KCNE1-related;  
KW Mink-related; long QT syndrome; cardiac arrhythmia;  
KW drug screening; knockout mouse; transgenic animal; ion channel disorder;  
KW fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
KW HERG; mutant; ds.  
XX  
OS Homo sapiens.  
XX Synthetic.  
XX  
PN M0200063434-A1.  
XX  
XX 26-OCT-2000.  
PF 14-APR-2000; 2000M0-0S10004.  
PR 15-APR-1999; 9908-0129404.  
XX (UTRN ) UNIV UTAH RES FOUND.  
XX (UTRA ) UNIV YALE.  
XX  
PI Abbott GM, Sesti F, Splawski I, Keating WT, Goldstein SM:  
XX WPI: 2000-672747/65.  
DR P-PSDB; AAB29593.  
XX  
PT Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for  
XX diagnosing and treating ion channel disorders, especially long QT  
XX syndrome -  
XX  
XX Claim 56; Page -; 132pp; English.

XX The invention relates to novel ion channel proteins related to  
XX KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
XX the invention are human and rat KCNE2 (MIRP1; AAB29585 and AAB29586,  
XX respectively); and human and mouse KCNE3 (MIRP2; AAB29588 and AAB29589,  
XX respectively). The cDNAs encoding these proteins are given in AAC64071-  
XX AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
XX potassium channels (I-Kr). Mutations in which are associated with long  
XX QT syndrome. The invention also relates to methods of diagnosing long QT  
XX syndrome using the KCNE2, KCNE3 or KCNE4 genes; a knockout mouse with a  
XX disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene; transgenic  
XX non-human animals expressing a transgenic animal comprising human KCNE2 and HERG  
XX DNA, and methods of and screening drugs for treating long QT syndrome  
XX using KCNE2 proteins (including mutants), nucleic acids encoding them  
XX and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
XX acids, and proteins may be used for diagnosing or treating ion channel  
XX disorders, especially long QT syndrome. Transgenic animals comprising  
XX KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
XX The invention also relates to methods of diagnosing long QT syndrome  
XX (MIRP1) specifically claimed for use in diagnostic and drug screening  
XX methods of the invention.  
XX Note: The present sequence is not shown in the specification, but is  
XX derived from the wild-type human KCNE2 cDNA sequence shown on page  
XX 118-119.  
XX  
XX Sequence 732 BP; 221 A; 151 C; 158 G; 202 T; 0 other:  
XX  
XX Query Match 99.6% Score 370.4; DB 21; Length 732;  
XX Best Local Similarity 99.7%; Prod. No. 6,7e-102;  
XX Matches 371; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

DB 1 ATGTCTATTATTCGCAATTTCACACAGACGCTGCTTCGGAAGATTTTAAAT 60  
|||||  
DB 74 ATGTCTATTATTCGCAATTTCACACAGACGCTGCTTCGGAAGATTTTAAAT 133

QY 61 ACTTATATGACAAATTTGGCCCGCCAGAACAGCTGACAAAGAGCCCTCCAGCAAA 120  
DB 134 ACTTATATGACAAATTTGGCCCGCCAGAACAGCTGACAAAGAGCCCTCCAGCAAA 193  
QY 121 GTTGATCTGAGAACTTCCTATATGTCATCGTGTACCTGAGATTTGGAATGTTTC 180  
|||||  
DB 194 GTTGATCTGAGAACTTCCTATATGTCATCGTGTACCTGAGATTTGGAATGTTTC 253  
QY 181 TCTTTCATCATCGTGTGACCATCTGCTGTGACCATCTGTGAATTCGAGAGCGGGAACATCC 240  
DB 254 TCTTTCATCATCGTGTGACCATCTGCTGTGACCATCTGTGAATTCGAGAGCGGGAACATCC 313  
QY 241 AATGACCCCTGACACAGTACATGTGACAGCATGTGACAGAGAAATTCGAGAGCGGGAACATCC 300  
DB 314 AATGACCCCTGACACAGTACATGTGACAGCATGTGACAGAGAAATTCGAGAGCGGGAACATCC 373  
QY 301 TTGATCTGAGAACTTCGAGAACTTCGAGAACTTCGAGAACTTCGAGAACTTCGAGAACTTC 360  
DB 374 TTGATCTGAGAACTTCGAGAACTTCGAGAACTTCGAGAACTTCGAGAACTTCGAGAACTTC 433  
QY 361 ATGTCCCTCTGGA 372  
|||||  
DB 434 ATGTCCCTCTGGA 445

RESULT 10  
AAC64084  
ID AAC64084 standard; DNA; 732 BP.  
XX  
AC AAC64084;  
XX  
XX 19-FEB-2001 (first entry)  
XX  
XX Human potassium channel protein KCNE2 (MIRP1) M54T mutant DNA.  
XX  
XX Human: KCNE2; MIRP1; potassium channel protein; KCNE1-related;  
XX Mink-related; long QT syndrome; cardiac arrhythmia;  
XX drug screening; knockout mouse; transgenic animal; ion channel  
XX fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
XX HERG; mutant; ds.  
XX  
XX Homo sapiens.  
XX Synthetic.  
XX  
XX M0200063434-A1.  
XX  
XX 26-OCT-2000.  
XX  
XX 14-APR-2000; 2000M0-0S10004.  
XX  
XX 15-APR-1999; 9908-0129404.  
XX (UTRA ) UNIV UTAH RES FOUND.  
XX (UTYA ) UNIV YALE.  
XX  
PI Abbott GM, Sesti F, Splawski I, Keating WT, Goldstein SM:  
XX WPI: 2000-672747/65.  
DR P-PSDB; AAB29594.  
XX  
PT Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for  
XX diagnosing and treating ion channel disorders, especially long QT  
XX syndrome -  
XX  
XX Claim 56; Page -; 132pp; English.

XX The invention relates to novel ion channel proteins related to  
XX KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
XX the invention are human and rat KCNE2 (MIRP1; AAB29585 and AAB29586,  
XX respectively); human and mouse KCNE3 (MIRP2; AAB29588 and AAB29589,  
XX respectively). The cDNAs encoding these proteins are given in AAC64071-  
XX AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
XX potassium channels (I-Kr). Mutations in which are associated with long  
XX QT syndrome. The invention also relates to methods of diagnosing long QT  
XX syndrome using the KCNE2, KCNE3 or KCNE4 genes; a knockout mouse with a  
XX disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene; transgenic  
XX non-human animals expressing a transgenic animal comprising human KCNE2 and HERG  
XX DNA, and methods of and screening drugs for treating long QT syndrome  
XX using KCNE2 proteins (including mutants), nucleic acids encoding them  
XX and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
XX acids, and proteins may be used for diagnosing or treating ion channel  
XX disorders, especially long QT syndrome. Transgenic animals comprising  
XX KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
XX The invention also relates to methods of diagnosing long QT syndrome  
XX (MIRP1) specifically claimed for use in diagnostic and drug screening  
XX methods of the invention.  
XX Note: The present sequence is not shown in the specification, but is  
XX derived from the wild-type human KCNE2 cDNA sequence shown on page  
XX 118-119.  
XX  
XX Sequence 732 BP; 221 A; 151 C; 158 G; 202 T; 0 other:  
XX  
XX Query Match 99.6% Score 370.4; DB 21; Length 732;  
XX Best Local Similarity 99.7%; Prod. No. 6,7e-102;  
XX Matches 371; Conservative 0; Mismatches 1; Indels 0; Gaps 0;



Db 314 AATGACCCCTACACGACATGTTGAGAGAGCTGGCAGGAAAGATACAGAGCCCAATTC 373  
 301 TTGAATCTAGAGAAATCGAAGGCCACACATCGATGAAACATTCGCTCGCGCTGCTTCAA 360  
 374 TTGAACTCTAGAGAAATCGAAGGCCACACATCGATGAAACATTCGCTCGCGCTTCAA 433  
 Oy 361 ATGATCCCTCGCA 372  
 434 ATGTCCCTCGCA 445  
 Db  
 RESULT 12  
 AAC64086 standard: DNA; 732 BP.  
 AAC64086;  
 AC AAC64086;  
 19-FEB-2001 (first entry)  
 D Human potassium channel protein KCNE2 (MIRP1) T9A mutant DNA.  
 D Human KCNE2; MIRP1; potassium channel protein; KCNE1-related;  
 K human-related; long QT syndrome; cardiac arrhythmia;  
 K drug screening; knockout mouse; transgenic animal; ion channel disorder;  
 K fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
 K HERG; mutant; ds.  
 K Homo sapiens.  
 K Synthetic.  
 OS  
 PN W0200063434-A1.  
 26-OCT-2000.  
 PD 14-APR-2000; 2000MO-US10004.  
 PR 15-APR-1999; 99US-0129404.  
 PA (UTAH ) UNIV UTAH RES FOUND.  
 PA (UYVA ) UNIV YALE.  
 PI Abbott GM, Seatl F, Splawski I, Reeling MF, Goldstein SAN;  
 PI WPI: 2000-672747/65.  
 DR P-PSDB: AAB29596.  
 XX  
 PT Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for  
 PT diagnosing and treating ion channel disorders, especially long QT  
 PT syndrome -  
 XX  
 XS Claim 56; Page -: 132pp; English.  
 CC The invention relates to novel ion channel proteins related to  
 CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
 CC the invention are human and rat KCNE2 (MIRP1; AAB29585 and AAB29586,  
 CC respectively); human and mouse KCNE3 (MIRP2; AAB29587 and AAB29588,  
 CC respectively); and human and mouse KCNE4 (MIRP3; AAB29589 and AAB29590,  
 CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
 CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
 CC channel (I<sub>Kr</sub>) in which are associated with long  
 CC QT syndrome. The invention also includes a transgenic animal comprising  
 CC a syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
 CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
 CC nonhuman animals comprising a heterologous ion channel protein gene  
 CC of the invention, and methods of and screening drugs for treating long QT syndrome  
 CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
 CC (including mutants), transgenic animals, and transgenic animals comprising  
 CC nucleic acids, and proteins may be used for diagnosing and treating ion channel  
 CC disorders, especially long QT syndrome. Transgenic animals comprising  
 CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
 CC The present sequence represents DNA encoding a mutant human KCNE2  
 CC (MIRP1) specifically claimed for use in diagnostic and drug screening

CC methods of the invention.  
 CC Note: The present sequence is not shown in the specification, but is  
 CC derived from the wild-type human KCNE2 cDNA sequence shown on page  
 CC 118-119.  
 CC  
 XX  
 S0 Sequence 732 BP; 220 A; 152 C; 158 G; 202 T; 0 other;  
 Query Match 99.6%; Score 370.4; DB 21; Length 732;  
 Best Local Similarity 99.7%; Pred. No. 6.7e-102;  
 Matches 371; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 Oy 1 ATGCTACTTATTCGAAATTTGACAGAGAGGCTGGAAGAGCGTCTCCGAGATTTTAT 60  
 Db 74 ATGCTACTTATTCGAAATTTGACAGAGAGGCTGGAAGAGCGTCTCCGAGATTTTAT 133  
 Oy 61 ACTTATATGACATATTCGCGCGAGAGAACACACGATGACGAGAGCGCTCCAGCCAA 120  
 Db 134 ACTTATATGACATATTCGCGCGAGAGAACACACGATGACGAGAGCGCTCCAGCCAA 193  
 Oy 121 GTTATGTCGACAGCTTCGACTATGATGATGATGATGATGATGATGATGATGATGAT 180  
 Db 194 GTTATGTCGACAGCTTCGACTATGATGATGATGATGATGATGATGATGATGATGAT 253  
 Oy 181 TCTTTCATCATGCTGCGCCACCTCGGTATGACACGTGGAATATCCAGAGAGCGGACACTC 240  
 Db 254 TCTTTCATCATGCTGCGCCACCTCGGTATGACACGTGGAATATCCAGAGAGCGGACACTC 313  
 Oy 241 AATGACCCCTACACGACATGTTGAGAGAGCTGGCAGGAAAGATACAGCCCAATTC 300  
 Db 314 AATGACCCCTACACGACATGTTGAGAGAGCTGGCAGGAAAGATACAGCCCAATTC 373  
 Oy 301 TTGAATCTAGAGAAATCGAAGGCCACACATCGATGAAACATTCGCTCGCGCTTCAA 360  
 Db 374 TTGAACTCTAGAGAAATCGAAGGCCACACATCGATGAAACATTCGCTCGCGCTTCAA 433  
 Oy 361 ATGATCCCTCGCA 372  
 434 ATGTCCCTCGCA 445  
 Db  
 RESULT 13  
 ABA4938 standard: DNA; 312 BP.  
 ABA4938;  
 01-FEB-2002 (first entry)  
 D Human breast cell single exon nucleic acid probe #8633.  
 D  
 D Human breast cell single exon probe: gene expression; breast;  
 D disease; cancer; ss.  
 D Homo sapiens.  
 OS  
 PN W02001517271-A2.  
 09-AUG-2001.  
 PD 30-JAN-2001; 2001MO-US000662.  
 PR 04-FEB-2000; 2000US-0180312.  
 PR 26-MAY-2000; 2000US-0207456.  
 PR 30-JUN-2000; 2000US-0608408.  
 PR 03-AUG-2000; 2000US-0632366.  
 PR 21-SEP-2000; 2000US-0234687.  
 PR 22-SEP-2000; 2000US-0236359.  
 PR 04-OCT-2000; 2000US-0024283.  
 PA (MOLE-) MOLECULAR DYNAMICS INC.  
 PI Penn SC, Hanzel DK, Chen W, Rank DB;

DR WPI: 2001-496933/54.  
 PR New spatially-addressable set of single exon nucleic acid probes,  
 PT specifically designed to amplify derived from human  
 PR breast, comprises number of single exon nucleic acid probes -  
 XX  
 PS Claim 4: SEQ ID NO 8633; 327bp + sequence listing; English.  
 CC  
 XX The invention relates to a spatially-addressable set of single exon  
 CC nucleic acid probes for measuring gene expression in a sample derived  
 CC from a specific tissue or organ. The probes involve contacting  
 CC the probes with a collection of detectably labeled cDNA  
 CC derived from mRNA of human breast, and then measuring the label  
 CC bound to each probe of the microarray. The probes are useful for  
 CC verifying the expression of regions of genomic DNA predicted to  
 CC encode proteins. They are useful for gene discovery, and for  
 CC determining predisposition and/or prognosing breast disease. Gene  
 CC expression analysis is useful for assessing the toxicity of chemical  
 CC agents and for the diagnosis of breast cancer. The microarray  
 CC diversity of probes for measuring gene expression, with far less bias  
 CC than expressed sequence tag microarrays. The method is suitable for  
 CC rapid production of functional information from genomic sequence. The  
 CC present sequence is a single exon nucleic acid probe of the invention.  
 CC Note: The sequence data for this patent did not form part of the  
 CC printed specification, but was obtained in electronic format directly  
 CC from Wipo at fcp.wipo.int/pub/linked\_Pct\_sequences.  
 XX  
 SQ Sequence 312 BP: 94 A; 73 C; 71 G; 74 T; 0 other;  
 Query Match 83.9%; Score 312; DB 22; Length 312;  
 Best Local Similarity 100.0%; Pred. No. 2e-84;  
 Matches 312; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Oy 54 TTTTATTCTTATATGAGCAATTTGGCGCGACAGCAACAGCTGAGAGGCCCTCCA 113  
 Db 1 TTTTATTCTTATATGAGCAATTTGGCGCGACAGCAACAGCTGAGAGGCCCTCCA 60  
 Oy 114 ACCCAAGTTCATGCTGAGCAATCTTCATCTGCTGCTGACCTGATGATGATG 173  
 Db 61 ACCCAAGTTCATGCTGAGCAATCTTCATCTGCTGCTGACCTGATGATGATG 120  
 Oy 174 AATGTCCTTTCATCATGTCGCGACCTGCTGACACCTGGAATTCAGAGCGGGA 233  
 Db 121 AATGTCCTTTCATCATGTCGCGACCTGCTGACACCTGGAATTCAGAGCGGGA 180  
 Oy 234 ACATGCAATGACCCCTACACCACTGATCTGATGAGCAATGCGAAGATGCAAG 293  
 Db 181 ACATGCAATGACCCCTACACCACTGATCTGATGAGCAATGCGAAGATGCAAG 240  
 Oy 294 CCAAACTTGATCTAGAGAAATCGAAGGCGACCACTGATGAGAAATTTGGTCGGTGG 353  
 Db 241 CCAAACTTGATCTAGAGAAATCGAAGGCGACCACTGATGAGAAATTTGGTCGGTGG 300  
 Oy 354 GTTCAAAATGTC 365  
 Db 301 GTTCAAAATGTC 312

RESULT 14  
 ABM67856 standard: DNM: 312 BP.  
 XX  
 AC ABM67856:  
 XX  
 DT 01-FEB-2002 (first entry)  
 XX  
 DE Human foetal liver single exon nucleic acid probe #16161.  
 XX  
 XX Human: foetal liver; gene expression, single exon nucleic acid probe. ss.  
 OS Homo sapiens.  
 XX  
 PN WO200157277-A2.

XX  
 XD 09-AUG-2001.  
 XX  
 XX 30-JAN-2001: 2001WO-US00669.  
 XX  
 PR 04-FEB-2000: 2000US-0180312.  
 PR 26-MAY-2000: 2000US-0207456.  
 PR 30-JUN-2000: 2000US-0608408.  
 PR 03-AUG-2000: 2000US-0632366.  
 PR 21-SEP-2000: 2000US-0234687.  
 PR 27-SEP-2000: 2000US-0236359.  
 PR 04-OCT-2000: 2000US-0024283.  
 XX  
 PA (MOLE-) MOLECULAR DYNAMICS INC.  
 PI Penn SG, Hanzel DK, Chen W, Rank DR.  
 DR WPI: 2001-483447/52.  
 PR Human genome-derived single exon nucleic acid probes useful for  
 PR analyzing gene expression in human fetal liver -  
 XX  
 PS Claim 4: SEQ ID NO 16161: 639bp + sequence listing; English.  
 CC  
 XX The invention relates to a single exon nucleic acid probe for  
 CC measuring human gene expression in a sample derived from human foetal  
 CC liver. The probes involve contacting the probes with a collection of  
 CC labeled cDNA derived from human foetal liver, and then measuring the  
 CC label bound to each probe of the microarray. The probes are useful for  
 CC verifying the expression of regions of genomic DNA predicted to  
 CC encode proteins. They are useful for gene discovery, and for  
 CC determining predisposition and/or prognosing breast disease. Gene  
 CC expression analysis is useful for assessing the toxicity of chemical  
 CC agents and for the diagnosis of breast cancer. The microarray  
 CC diversity of probes for measuring gene expression, with far less bias  
 CC than expressed sequence tag microarrays. The method is suitable for  
 CC rapid production of functional information from genomic sequence. The  
 CC present sequence is a single exon nucleic acid probe of the invention.  
 CC Note: The sequence data for this patent did not form part of the  
 CC printed specification, but was obtained in electronic format directly  
 CC from Wipo at fcp.wipo.int/pub/linked\_Pct\_sequences.  
 XX  
 SQ Sequence 312 BP: 94 A; 73 C; 71 G; 74 T; 0 other;  
 Query Match 83.9%; Score 312; DB 22; Length 312;  
 Best Local Similarity 100.0%; Pred. No. 2e-84;  
 Matches 312; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Oy 54 TTTTATTCTTATATGAGCAATTTGGCGCGACAGCAACAGCTGAGAGGCCCTCCA 113  
 Db 1 TTTTATTCTTATATGAGCAATTTGGCGCGACAGCAACAGCTGAGAGGCCCTCCA 60  
 Oy 114 ACCCAAGTTCATGCTGAGCAATCTTCATCTGCTGCTGACCTGATGATGATG 173  
 Db 61 ACCCAAGTTCATGCTGAGCAATCTTCATCTGCTGCTGACCTGATGATGATG 120  
 Oy 174 AATGTCCTTTCATCATGTCGCGACCTGCTGACACCTGGAATTCAGAGCGGGA 233  
 Db 121 AATGTCCTTTCATCATGTCGCGACCTGCTGACACCTGGAATTCAGAGCGGGA 180  
 Oy 234 ACATGCAATGACCCCTACACCACTGATCTGATGAGCAATGCGAAGATGCAAG 293  
 Db 181 ACATGCAATGACCCCTACACCACTGATCTGATGAGCAATGCGAAGATGCAAG 240  
 Oy 294 CCAAACTTGATCTAGAGAAATCGAAGGCGACCACTGATGAGAAATTTGGTCGGTGG 353  
 Db 241 CCAAACTTGATCTAGAGAAATCGAAGGCGACCACTGATGAGAAATTTGGTCGGTGG 300  
 Oy 354 GTTCAAAATGTC 365  
 Db 301 GTTCAAAATGTC 312

RESULT 15  
 ABA34913 standard: DNM: 312 BP.  
 XX  
 AC ABA34913:  
 XX  
 DT 23-JAN-2002 (first entry)  
 XX

DE Probe #13379 for gene expression analysis in human heart cell sample.  
 XX Human: gene expression: heart; microarray; vascular system; probe;  
 XX cardiovascular disease; hypertension; cardiac arrhythmias;  
 XX congenital heart disease; 5q.  
 XX Homo sapiens.  
 XX MO200157274-A2.  
 XX  
 XX 09-AUG-2001.  
 XX  
 XX 30-JAN-2001; 2001MO-US00666.  
 XX  
 XX 04-FEB-2000; 2000US-0180312.  
 XX 26-MAY-2000; 2000US-0207456.  
 XX 30-JUN-2000; 2000US-0608408.  
 XX 03-AUG-2000; 2000US-0632386.  
 XX 21-SEP-2000; 2000US-0234687.  
 XX 27-SEP-2000; 2000US-0234587.  
 XX 04-OCT-2000; 2000US-0024263.  
 XX  
 XX (MOLE-) MOLECULAR DYNAMICS INC.  
 XX  
 XX Penn SC, Hanzel DK, Chen W, Rank DR;  
 XX WPI; 2001-488899/53.  
 XX  
 XX Single exon nucleic acid probes for analyzing gene expression in human  
 PT hearts -  
 PS  
 PS Claim 4: SEQ ID NO 13379; 530bp; English.  
 XX  
 XX The present invention relates to single exon nucleic acid probes for  
 XX analyzing gene expression in a sample derived from human heart. The  
 XX present sequence from the probes published for the purpose of  
 CC predicting, measuring and displaying gene expression in samples derived  
 CC from the human heart via microarrays. By measuring gene expression, the  
 CC probes are useful for predicting, diagnosing, grading, staging,  
 CC monitoring and prognosing diseases of the human heart and vascular system  
 CC e.g. cardiovascular disease, hypertension, cardiac arrhythmias and  
 CC congenital heart disease.  
 CC The present state of the art for this patent did not form part of the printed  
 CC specification but was obtained in electronic format directly from WPI  
 CC at ftp.wpi.int/pub/published\_pat\_sequences.  
 XX  
 XX Sequence 312 BP; 94 A; 73 C; 71 G; 74 T; 0 other;  
 XX  
 XX Query Match 83.9%; Score 312; DB 22; Length 312;  
 XX Best Local Similarity 100.0%; Pred. No. 26-84;  
 XX Matches 312; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
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 XX 54 TTTTATCTATTATGAGCATTTGGCGGACAGACACAGCATGACAGAGGCGCTTCCA 113  
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 XX 1 TTTTATCTATTATGAGCATTTGGCGGACAGACACAGCATGACAGAGGCGCTTCCA 60  
 XX  
 XX 114 AGCCAAAGTAGAGCTGAGAACTTCATGATGTCATCTCTGACCTCATGATGATG 173  
 XX |  
 XX 61 AGCCAAAGTAGAGCTGAGAACTTCATGATGTCATCTCTGACCTCATGATGATG 120  
 XX  
 XX 174 ATGTGTCCTTTTCATCTGTCGCGCATCTGCTGAGACACTGTGAATCCAGAGACGGA 233  
 XX |  
 XX 121 ATGTGTCCTTTTCATCTGTCGCGCATCTGCTGAGACACTGTGAATCCAGAGACGGA 180  
 XX  
 XX 234 AACTGCATAGACCCCTACACACAGTACATTGTGAGGACTGCGAGAGAAAGTACAGAG 293  
 XX |  
 XX 181 AACTGCATAGACCCCTACACACAGTACATTGTGAGGAGCTGCGAGAGAAAGTACAGAG 240  
 XX  
 XX 294 CCAAACTTGATCTGAGAAAGATCCAGACCATCATGATGAGAACATTTGGTGGCGTGG 353  
 XX |  
 XX 241 CCAAACTTGATCTGAGAAAGATCCAGACCATCATGATGAGAACATTTGGTGGCGTGG 300  
 XX  
 XX 354 GTTCAAAATATGTC 365

DB ||||| 312  
 301 GTTCAAAATATGTC 312  
 Search completed: October 24, 2002, 13:19:51  
 Job time : 201 secs





GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: October 24, 2002, 13:16:15 : Search time 43 seconds  
(without alignments)

2125.015 Million cell updates/sec

US-09-550-163-1\_COPY\_74\_445

Sequence score: 372  
Sequence: 1 agtgcactatcaccatt.....gttcaaaagccccccgga 372

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 383533 seqs, 122816752 residues

Total number of hits satisfying chosen parameters: 767066

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 08  
Maximum Match 1008  
Listing first 45 summaries

Database :

Issued Patents NA.\*  
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2: /cgn2.6/prodata/2/ina/5B.COMB.seq.\*  
3: /cgn2.6/prodata/2/ina/6A.COMB.seq.\*  
4: /cgn2.6/prodata/2/ina/6B.COMB.seq.\*  
5: /cgn2.6/prodata/2/ina/PCFOS.COMB.seq.\*  
6: /cgn2.6/prodata/2/ina/backlist1.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	53.2	14.3	398	1	US-08-118-101A-5
2	53.2	14.3	1703	3	US-09-135-021-77
3	53.2	14.3	1703	4	US-09-135-020-3
4	53.2	14.3	1703	4	US-09-135-010A-3
5	36.6	9.7	1382	1	US-08-110-262A-1
6	33.6	9.0	665	5	US-09-059-896-2
7	33.6	8.9	606	4	US-09-328-111-133
8	33.6	8.9	606	4	US-09-307-143-3
9	33.6	8.9	2912	4	US-08-998-416-303
10	32.4	8.7	837	4	US-09-426-998-3
11	30.6	8.2	6822	4	US-08-445-886-5
12	29.8	8.2	2264	4	US-09-726-399A-18
13	29.8	8.0	246240	2	US-08-724-394A-20
14	29.8	8.0	246240	2	US-08-724-394A-21
15	29.8	8.0	246240	2	US-08-724-394A-22
16	29.8	8.0	246240	2	US-08-724-394A-23
17	29.6	8.0	462	3	US-08-863-813A-33
18	29.6	8.0	3273	6	5516630-1
19	29.6	8.0	7486	3	US-08-445-886-5
20	29.6	8.0	7486	3	US-08-445-886-5
21	29.6	8.0	7486	3	US-08-445-886-5
22	29.6	8.0	7493	3	US-08-445-886-1
23	29.6	8.0	7493	3	US-08-397-233-1
24	29.6	8.0	7493	3	US-09-111-387-1
25	29.4	7.9	3947	4	US-08-915-762-47
26	29.4	7.9	3947	4	US-09-285-028-47
27	29.4	7.9	3947	4	US-09-106-582-47

c 28	29.2	7.8	352	4	US-09-439-313-421	Sequence 421, App
c 29	29.2	7.8	630	4	US-08-646-695-13	Sequence 13, App1
c 30	29.2	7.8	630	4	US-08-646-695-14	Sequence 14, App1
c 31	29.2	7.8	630	5	PCF-US96-06053-13	Sequence 13, App1
c 32	29.2	7.8	230	4	PCF-US95-06053-14	Sequence 14, App1
c 33	29.2	7.8	14311	4	US-08-646-695-1	Sequence 1, App1
c 34	29.2	7.8	14311	4	US-08-646-695-1	Sequence 1, App1
c 35	29.2	7.8	14311	4	US-08-646-695-7	Sequence 7, App1
c 36	29.2	7.8	14311	5	PCF-US96-06053-7	Sequence 7, App1
c 37	29.2	7.8	14311	5	PCF-US96-06053-7	Sequence 7, App1
c 38	29	7.8	744	3	US-08-969-644-17	Sequence 17, App1
c 39	29	7.8	744	3	US-08-444-189-17	Sequence 17, App1
c 40	29	7.8	1964	4	US-08-444-189-17	Sequence 17, App1
c 41	29	7.8	7502	3	US-08-969-644-6	Sequence 6, App1
c 42	29	7.8	7502	3	US-08-444-189-6	Sequence 6, App1
c 43	29	7.8	7502	3	US-08-444-189-6	Sequence 6, App1
c 44	29	7.8	7502	4	US-08-468-544-6	Sequence 6, App1
c 45	28.8	7.7	2168	2	US-08-633-879C-1	Sequence 1, App1

## ALIGNMENTS

RESULT 1  
US-08-118-101A-5  
; Sequence 5, App10818101A  
; Patent No. 5620892  
; INVENTOR: Stephen E.  
; GENERAL INFORMATION:  
; APPLICANT: Knickerbocker, Aron M.  
; TITLE OF INVENTION: A STRAIN OF SACHAROMYCES CEREVISIAE  
; NUMBER OF SEQUENCES: 16  
; CORRESPONDENCE ADDRESS:  
; ADDRESS: Burton Rodney  
; CITY: Princeton  
; STATE: New Jersey  
; COUNTRY: U.S.A.  
; ZIP: 08543-4000  
; COMPUTER READABLE FORM:  
; MODUL TYPE: floppy disk  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/118-101A  
; FILING DATE:  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; REGISTRATION NUMBER: 33,111  
; REFERENCE/DOCKET NUMBER: DC27  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (609) 252-5901  
; INFORMATION FOR SEQ ID NO: 5:  
; SOURCE: CHLOROPHYLLS  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; FEATURE:  
; NAME/KEY: CDS  
; VALUE: 1..398  
US-08-118-101A-5

Query Match 14.3% Score 53.2: DB 1: Length 398:  
Best Local Similarity 63.6% Pred. No. 1.3e+06:  
Matches 98: Conservative 0: Mismatches 53: Indels 3: Gaps 1:

OY 150 CCTGTACCTCATGTGATGGATGGATGTTCTTTCATCATCATGTGGCCATCTCTGTGAG 209  
 DB 141 CCTGTACCTCATGTGATGGATGGATGTTCTTTCATCATCATGTGGCCATCTCTGTGAG 200  
 OY 210 CACGTGGAAATCCAGAGGAGGAAACCTCCATCCAGCCCTCCAGCATCATGTTG---T 266  
 DB 201 CTAATCCGCTCCAGAGGAGGAGGAAACCTCCATCCAGCCCTCCAGCATCATGTTG 260  
 OY 267 AGAGAGCTGGCAGAGAAAGTACAAAGGCCAATTC 300  
 DB 261 CCAATCCGCTCCAGAGGAGGAGGAAACCTCTATCTC 294

## RESULT 2

US-09-135-021-77  
 Sequence 77, Application US/09135021A

Patent No. 6150104  
 GENERAL INFORMATION:  
 APPLICANT: Spilawski, Igor  
 TITLE OF INVENTION: A HOMOZYGUS MUTATION IN KYLOT1 WHICH CAUSES JERVELL  
 FILE REFERENCE: 2323-128  
 CURRENT APPLICATION NUMBER: 05/09/135, 021A  
 EARLIER APPLICATION NUMBER: 05/09/135, 021A  
 EARLIER FILING DATE: 1997-06-13  
 EARLIER APPLICATION NUMBER: 60/094,475  
 EARLIER FILING DATE: 1997-06-13  
 EARLIER APPLICATION NUMBER: 60/094,477  
 EARLIER FILING DATE: 1998-07-29  
 NUMBER OF SEQ ID NOS: 80  
 SOFTWARE: PatentIn Ver. 2.0  
 SEQ ID NO 1703  
 TYPE: DNA  
 ORGANISM: Homo sapiens  
 FEATURE:  
 NAME/KEY: CDS  
 LOCATION: (193)..(579)  
 US-09-135-021-77

## Query Match

Best Local Similarity 63.6%; Pred. No. 3e-08; Length 1703;  
 Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;

OY 150 CCTGTACCTCATGTGATGGATGGATGTTCTTTCATCATCATGTGGCCATCTCTGTGAG 209  
 DB 324 CCTGTACCTCATGTGATGGATGGATGTTCTTTCATCATCATGTGGCCATCTCTGTGAG 200  
 OY 210 CACGTGGAAATCCAGAGGAGGAAACCTCCATCCAGCCCTCCAGCATCATGTTG---T 266  
 DB 384 CTAATCCGCTCCAGAGGAGGAGGAAACCTCCATCCAGCCCTCCAGCATCATGTTG 260  
 OY 267 AGAGAGCTGGCAGAGAAAGTACAAAGGCCAATTC 300  
 DB 444 CCAATCCGCTCCAGAGGAGGAGGAAACCTCTATCTC 294

## RESULT 3

US-09-135-020-3  
 Sequence 3, Application US/09135020

Patent No. 6274332  
 GENERAL INFORMATION:  
 APPLICANT: Spilawski, Mark T.  
 APPLICANT: Spilawski, Igor  
 APPLICANT: Spilawski, Michael C.  
 TITLE OF INVENTION: MUTATIONS IN THE KONE1 GENE ENCODING HUMAN MIK WHICH  
 TITLE OF INVENTION: CAUSE ARRYTHMIA SUSCEPTIBILITY THEREBY ESTABLISHING  
 FILE REFERENCE: 2323-131  
 CURRENT APPLICATION NUMBER: US/09/135, 020  
 EARLIER APPLICATION NUMBER: 08/921,068  
 EARLIER FILING DATE: 1997-08-29

EARLIER APPLICATION NUMBER: 08/739,383  
 EARLIER FILING DATE: 1996-10-29  
 EARLIER APPLICATION NUMBER: 60/019,014  
 EARLIER FILING DATE: 1996-10-29  
 EARLIER APPLICATION NUMBER: 60/094,477  
 EARLIER FILING DATE: 1998-07-29  
 NUMBER OF SEQ ID NOS: 114  
 SOFTWARE: PatentIn Ver. 2.0  
 SEQ ID NO 3  
 TYPE: DNA  
 ORGANISM: Homo sapiens  
 FEATURE:  
 NAME/KEY: CDS  
 LOCATION: (193)..(579)  
 US-09-135-020-3

## Query Match

Best Local Similarity 63.6%; Pred. No. 3e-08; Length 1703;  
 Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;

OY 150 CCTGTACCTCATGTGATGGATGGATGTTCTTTCATCATCATGTGGCCATCTCTGTGAG 209  
 DB 324 CCTGTACCTCATGTGATGGATGGATGTTCTTTCATCATCATGTGGCCATCTCTGTGAG 200  
 OY 210 CACGTGGAAATCCAGAGGAGGAAACCTCCATCCAGCCCTCCAGCATCATGTTG---T 266  
 DB 384 CTAATCCGCTCCAGAGGAGGAGGAAACCTCCATCCAGCCCTCCAGCATCATGTTG 260  
 OY 267 AGAGAGCTGGCAGAGAAAGTACAAAGGCCAATTC 300  
 DB 444 CCAATCCGCTCCAGAGGAGGAGGAAACCTCTATCTC 294

## RESULT 4

US-09-135-010A-3  
 Sequence 3, Application US/09135010A

Patent No. 6277978  
 GENERAL INFORMATION:  
 APPLICANT: Keating, Mark T.  
 APPLICANT: Keating, Mark T.  
 APPLICANT: Curran, Mark R.  
 APPLICANT: Landers, Gregory M.  
 APPLICANT: Connors, Timothy D.  
 APPLICANT: Burn, Timothy C.  
 APPLICANT: Spilawski, Igor  
 TITLE OF INVENTION: KYLOT1 - A LONG QT SYNDROME GENE  
 FILE REFERENCE: 2323-133  
 CURRENT APPLICATION NUMBER: US/09/135, 010A  
 EARLIER APPLICATION NUMBER: 08/921,068  
 EARLIER FILING DATE: 1998-08-17  
 EARLIER APPLICATION NUMBER: 60/094,477  
 EARLIER FILING DATE: 1998-07-29  
 EARLIER APPLICATION NUMBER: 08/921,068  
 EARLIER FILING DATE: 1997-08-29  
 EARLIER APPLICATION NUMBER: 05/739,383  
 EARLIER FILING DATE: 1996-10-29  
 EARLIER APPLICATION NUMBER: 60/019,014  
 EARLIER FILING DATE: 1995-12-22  
 NUMBER OF SEQ ID NOS: 116  
 SOFTWARE: PatentIn Ver. 2.0  
 SEQ ID NO 3  
 TYPE: DNA  
 ORGANISM: Homo sapiens  
 FEATURE:  
 NAME/KEY: CDS  
 LOCATION: (193)..(579)  
 US-09-135-010A-3

Query Match  
 Best Local Similarity 63.6%; Pred. No. 3e-08; Length 1703;  
 Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;



QY 115 GCGAAGTGAATGCTGAGAACCTTACATGATGCTGATACCTGATGCTGATGATGA 174  
 DB 862 GCGAAGGCTGCTGGGTGTCAGACGATCATCTGACAGGCGCCATGATCTGCTG 921  
 QY 175 ATGTCTCTTTCATCATCTGAGCCATCTGCTGAGCACTGTGAATCCAGAGCGGGA 234  
 DB 922 CTGATCAATTCATCTCTCTTTCATGATGCTGATGCTGATGCTGATGCTGATG 981  
 QY 235 GATCTGATGATGACCCCTGATGATGATGATGATGATGATGATGATGATGATGAT 262  
 DB 982 TCCACACGCTGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 1009  
  
 RESULT 7  
 US-09-069-896-2  
 Sequence 2, Application US/09069896  
 GENERAL INFORMATION:  
 APPLICANT: Hillman, Jennifer L.  
 APPLICANT: Peterson, Chandra  
 APPLICANT: Corley, Neil C.  
 TITLE OF INVENTION: DELAYED RECTIFIER POTASSIUM  
 TITLE OF INVENTION: CHANNEL HOMOLOG  
 ADDRESS: SPOKANE, ID 83400-4000  
 ADDRESS: INCYTE Pharmaceuticals, Inc.  
 STREET: 3174 Porter Drive  
 CITY: Palo Alto  
 STATE: CA  
 COUNTRY: USA  
 COMPUTER NAME: 3174-1000  
 MEDIUM TYPE: Diskette  
 OPERATING SYSTEM: IBM Compatible  
 SOFTWARE: FASTEDS for Windows Version 2.0  
 CURRENT APPLICATION DATA:  
 FILING DATE: NUMBER: US/09/069,896  
 CLASSIFICATION:  
 PRIOR APPLICATION NUMBER:  
 FILING DATE:  
 ATTORNEY/AGENT INFORMATION:  
 NAME: Gertrone, Michael C.  
 REFERENCE/DOCKET NUMBER: PF-0507 US  
 TELEPHONE: 650-855-0555  
 TELEFAX: 650-845-4166  
 TELEX:  
 INFORMATION FOR SEQ ID NO: 2:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 913  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 IMMEDIATE SOURCE:  
 LIBRARY: BRSTN073  
 CLONE: 637471  
 US-09-069-896-2  
  
 Query Match 9.08; Score 33.6; DB 3; Length 645;  
 Best Local Similarity 53.98; Pred. No. 0.081; Indels 0; Gaps 0;  
 Matches 69; Conservative 0; Mismatches 59;

QY 270 GCACTGGC 277  
 DB 320 GGAGCGGC 327  
  
 RESULT 8  
 US-09-328-111-133  
 Sequence 133, Application US/09328111  
 GENERAL INFORMATION:  
 APPLICANT: Enders, Wilson O.  
 APPLICANT: Steinmann, Kathleen E.  
 APPLICANT: Astle, Jon H.  
 APPLICANT: Burgess, Christopher C.  
 APPLICANT: Bushnell, Steven E.  
 APPLICANT: Carroll III, Eddie  
 APPLICANT: Dettlo, Theodore J.  
 APPLICANT: Ford, Donna M.  
 APPLICANT: Lewis, Marcia E.  
 APPLICANT: Monahan, John E.  
 APPLICANT: Schlegel, Robert  
 TITLE OF INVENTION: NOVEL HUMAN GENES AND GENE EXPRESSION  
 TITLE OF INVENTION: PRODUCTS  
 CURRENT APPLICATION NUMBER: US/09/328,111  
 CURRENT FILING DATE: 1999-06-08  
 EARLIER APPLICATION NUMBER: US 60/088,801  
 EARLIER FILING DATE: 1998-06-10  
 NUMBER OF SEQ ID NOS: 850  
 SOFTWARE: FASTEDS for Windows Version 3.0  
 SEQ ID NOS: 1-850  
 LENGTH: 606  
 TYPE: DNA  
 ORGANISM: Homo sapiens  
 FEATURE:  
 NAME/KEY: misc\_feature  
 LOCATION: (1)..(606)  
 OTHER INFORMATION: n = A,T,C or G  
 US-09-328-111-133  
  
 Query Match 8.98; Score 33; DB 4; Length 606;  
 Best Local Similarity 58.88; Pred. No. 0.13;  
 Matches 57; Conservative 0; Mismatches 40; Indels 0; Gaps 0;  
 QY 135 CTCTTCACATGATCTGCTACCTGATGATGATGATGATGATGATGATGATGATGAT 194  
 DB 124 CTCTTCCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 103  
 QY 195 GCGCATCTGCTGAGCAGCTGTGAATCCAGAGCGG 231  
 DB 184 TCTCCAGAAAAGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 220  
  
 RESULT 9  
 US-09-307-143-3/C  
 Sequence 3, Application US/09307143  
 Patent No. 6335157  
 GENERAL INFORMATION:  
 APPLICANT: Gonzalez C.  
 APPLICANT: Lange, B.  
 TITLE OF INVENTION: METHODS BASED ON LOCALIZATION OF HSP90 TO THE  
 TITLE OF INVENTION: HSP90 SOMA  
 FILE REFERENCE: 9882-003  
 CURRENT APPLICATION NUMBER: US/09/307,143  
 CURRENT FILING DATE: 1999-05-07  
 NUMBER OF SEQ ID NOS: 12  
 SOFTWARE: PatentIn Ver. 2.1  
 SEQ ID NOS: 1-12  
 LENGTH: 2782  
 TYPE: DNA  
 ORGANISM: Homo sapiens





```

? APPLICANT: Feder, John N.
? APPLICANT: Krommal, Gregory S.
? APPLICANT: Lauer, Peter M.
? APPLICANT: Ruddy, David A.
? APPLICANT: Thomas Winston
? APPLICANT: Thompson, J. Kentia
? APPLICANT: Wolff, Roger K.
? TITLE OF INVENTION: Megabase Transcript Map: No. 5872237el
? NUMBER OF SEQUENCES: 31
? CORRESPONDENCE ADDRESS:
? ADDRESSER: TOWNSEND and TOWNSEND and CREW LLP
? STREET: 100 Montecarlo Center, 8th Floor
? CITY: San Francisco
? STATE: CA
? COUNTRY: USA
? ZIP: 94111-3834
? COMPUTER READABLE FORM:
? MEDIUM TYPE: Floppy disk
? COMPUTER: IBM PC compatible
? SOFTWARE: IBM DOS/MS-DOS
? SOFTWARE: PatentIn Release #1.0, Version #1.30
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: US/08/724,394A
? FILING DATE: 01-OCT-1996
? CLASSIFICATION: 536
? ATTORNEY/AGENT INFORMATION:
? NAME: Fitts, Renee A. 35,136
? REFERENCE NUMBER: 017957-000100
? TELECOMMUNICATION INFORMATION:
? TELEPHONE: 415-576-0200
? TELEFAX: 415-576-0300
? INFORMATION FOR SEQ ID NO: 21:
? SEQUENCE CHARACTERISTICS:
? LENGTH: 246240 base pairs
? TYPE: nucleic acid
? STRANDEDNESS: not relevant
? TOPOLOGY: not relevant
? MOLECULE TYPE: cDNA
? FEATURE:
? NAME/KEY: misc_feature
? POSITIVE: 1..246240
? OTHER INFORMATION: /note="HLA-H-CONTIG"
US-08-724-394A-21

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Query Match 8.0%; Score 29.8; DB 2; Length 246240;

Best Local Similarity 60.5%; Pred. No. 50;

Matches 49; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

OY 152 TGTACCTCAGTCGATGATGTGAGATGTCCTTTCATCATCGTCGCGCATCTGGTGAGA 211

DB 29016 TGTTCACCATGCGCATGATCGCATGACACTCAGCGCTGGTATTCATGCTGTAAGT 29075

OY 212 CTGTGAATCCAGAGACGGG 232

DB 29076 TTGAGAGTCATGATPAGAG 29096

Search completed: October 24, 2002, 14:20:05  
Job time : 147 secs





GenCore version 5.1.3  
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 24, 2002, 13:15:40 : Search time 1605 Seconds

(without alignments)  
3128.266 Million cell updates/sec

File: US-09-550-163-1\_COPY\_74\_445

Reflect score: 372  
1 atgcctactatcaatt.....gtccaaatgcccccca 372

Scoring table: IDENTITY\_NUC  
Gapop 10.0, Gapext 1.0

Searched: 13736207 seqs, 674847542 residues

Total number of hits satisfying chosen parameters: 27472414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

ESR:\*

1: em\_gstha:\*\*\*  
2: em\_gstlm:\*\*\*  
3: em\_gstlm:\*\*\*  
4: em\_gstlm:\*\*\*  
5: em\_gstlm:\*\*\*  
6: em\_gstlm:\*\*\*  
7: em\_gstlm:\*\*\*  
8: em\_gstlm:\*\*\*  
9: gb\_aatl:\*\*\*  
10: gb\_aatl:\*\*\*  
11: gb\_aatl:\*\*\*  
12: gb\_gst:\*\*\*  
13: gb\_gst:\*\*\*  
14: gb\_gst:\*\*\*  
15: em\_gstlm:\*\*\*  
16: em\_gstlm:\*\*\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the total being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
c 1	357.8	96.2	803	10	BG208163
2	336	90.3	410	9	A1962650
3	336	90.3	410	9	A1962650
4	331.8	89.2	399	9	A1962650
5	328	88.2	372	9	A1246239
6	269.6	72.5	1691	11	AK008619
7	260.8	70.1	470	10	DB5797
8	245	65.9	1003	10	BG261965
9	177.8	47.8	746	10	BG21966
10	168.2	45.2	751	10	BG334225
11	168.2	45.2	751	10	BG334225
12	119.4	32.1	9	AM869303	
13	99	26.6	311	9	AM869303
c 14	90	24.5	272	9	BB564873
15	57.6	15.5	272	9	BB564873
16	57.6	15.4	358	10	BB486735
17	53.2	14.3	716	10	BB459541
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					A1962650 A048012
					A1962650 A048012
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					A1246239 A1246239
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					A1246239 A1246239

RESULT 1					
BG208163/c	803 bp	mRNA	linear	EST 21-APR-2001	
LOCUSTATION					
ACCESSION					
VERSION					
KEYWORDS					
SOURCE					
ORGANISM					
human.					
Homo sapiens					
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;					
Mammalia; Eutheria; Primates; Carnivora; Homnidae; Homo.					
REFERENCE					
AUTHORS					
Harrington,J., Sherf, B., Rindlett, S., Jackson, P.D., Perry, R.,					
1 (bases 1 to 803)					
Levine, L., Goodman, D., Thompson, R., Thompson, J.					
Farmer, I., Goodman, D., McEligott, K., Roovers, S., Myers, R., Smith					
, B., Veloso, N., Kille, A., Hess, J., Colburn, K., Lo, K., Offenbacher					
, J., Danzaj, J., and Duerck, M.					
Creation of genome-wide protein expression libraries using random					
activation of gene expression					
Nat. Biotechnol. 19 (5), 440-445 (2001)					
TITLE					
JOURNAL					
MEDLINE					
COMMENT					
Contact: Scott J. Cain					
3200					
Atkarsky, Inc. Ave, Cleveland, OH 44115, USA					
Tel: 216 431 9900					
Fax: 216 361 9596					
Email: scalc@atkarsky.com					
High quality sequence stop: 550.					
Location/Qualifiers					
1..803					
/organism="Homo sapiens"					
/db_xref="taxon:9606"					
/clone-lib="Atkarsky RAGE Library"					
/note="Use: Creation of Genome-wide Protein Expression					
libraries using Random Activation of Gene Expression"					
Nature Biotechnology, In press. Note that even though the					
cell type indicated is H1080, since a random activation					
method was used, these sequence tags are not necessarily					
expressed in H1080 under normal circumstances."					
FEATURES					
source					

## ALIGNMENTS

RESULT 1					
BG208163/c	803 bp	mRNA	linear	EST 21-APR-2001	
LOCUSTATION					
ACCESSION					
VERSION					
KEYWORDS					
SOURCE					
ORGANISM					
human.					
Homo sapiens					
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;					
Mammalia; Eutheria; Primates; Carnivora; Homnidae; Homo.					
REFERENCE					
AUTHORS					
Harrington,J., Sherf, B., Rindlett, S., Jackson, P.D., Perry, R.,					
1 (bases 1 to 803)					
Levine, L., Goodman, D., Thompson, R., Thompson, J.					
Farmer, I., Goodman, D., McEligott, K., Roovers, S., Myers, R., Smith					
, B., Veloso, N., Kille, A., Hess, J., Colburn, K., Lo, K., Offenbacher					
, J., Danzaj, J., and Duerck, M.					
Creation of genome-wide protein expression libraries using random					
activation of gene expression					
Nat. Biotechnol. 19 (5), 440-445 (2001)					
TITLE					
JOURNAL					
MEDLINE					
COMMENT					
Contact: Scott J. Cain					
3200					
Atkarsys, Inc. Ave, Cleveland, OH 44115, USA					
Tel: 216 431 9900					
Fax: 216 361 9596					
Email: scalc@atkarsys.com					
High quality sequence stop: 550.					
Location/Qualifiers					
1..803					
/organism="Homo sapiens"					
/db_xref="taxon:9606"					
/clone-lib="Atkarsys RAGE Library"					
/note="Use of Creation of Genome-wide Protein Expression					
libraries using Random Activation of Gene Expression"					
Nature Biotechnology, In press. Note that even though the					
cell type indicated is H1080, since a random activation					
method was used, these sequence tags are not necessarily					
expressed in H1080 under normal circumstances."					

BASE COUNT 222 a 172 c 171 g 238 t  
ORIGIN  
Query Match 96.2%; Score 357.8; DB 10; Length 803;  
Best Local Similarity 99.2%; Pred. No. 1.2e-96;  
Matches 370; Conservative 0; Mismatches 2; Indels 1; Gaps 1;  
Oy 1 ATGCTTACTTATTCATTCACAGAGAGCTGGAGAGAGCTCTCCG-AAAGATTATTTAT 59  
Db 668 TACCTTATATGCAATATGCGCCAGCAACAGCCGTGAGAGAGAGAGAGAGAGAGCA 629  
Oy 60 TACCTTATATGCAATATGCGCCAGCAACAGCCGTGAGAGAGAGAGAGAGAGAGCA 629  
Db 628 TACCTTATATGCAATATGCGCCAGCAACAGCCGTGAGAGAGAGAGAGAGAGAGCA 629  
Oy 120 ACTGATATGTCAGAACCTTCTACTATATGTCATCTCTCTCTCTCTCTCTCTCTCT 179  
Db 568 AGTATATATGTCAGAACCTTCTACTATATGTCATCTCTCTCTCTCTCTCTCTCTCT 179  
Oy 180 CTCTTTCATCATCTGTCGACCTCTGTCGACCTCTGTCGACCTCTGTCGACCTCTG 239  
Db 508 CTCTTTCATCATCTGTCGACCTCTGTCGACCTCTGTCGACCTCTGTCGACCTCTG 239  
Oy 240 CATTGACCCCTGACCAACGATGATGATGATGATGATGATGATGATGATGATGATG 299  
Db 448 CATTGACCCCTGACCAACGATGATGATGATGATGATGATGATGATGATGATGATG 299  
Oy 300 CTGATATCTAGAGAAATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 359  
Db 388 CTGATATCTAGAGAAATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 359  
Oy 360 AATGTCCTGCTGCA 372  
Db 328 AATGTCCTGCTGCA 316

RESULT 2  
A1962650 410 bp mRNA linear EST 09-MAR-2000  
w412603.x1 NCI CGAP GC6 Homo sapiens RNA clone IMAGE:230895.3  
DEFINITION  
SIMILAR TO SW:MIK HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM  
CHANNEL PROTEIN ; mRNA sequence.  
ACCESSION  
A1962650.1 GI:5755363  
VERSION  
A1962650  
KEYWORDS  
EST  
ORGANISM  
Homo sapiens  
Human  
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.  
REFERENCE  
NCI CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.  
1 (bases 1 to 410)  
AUTHORS  
National Cancer Institute, Cancer Genome Anatomy Project (CGAP).  
TITLE  
Tumor Gene Index  
JOURNAL  
Contact: Robert Strausberg, Ph.D.  
COMMENT  
Email: [scap@f-mail.nih.gov](mailto:scap@f-mail.nih.gov)  
R. Emmert-Buck, M.D., Ph.D.  
CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima  
Bonaldo, Ph.D.  
DNA Sequencing by: Greg Lennon, Ph.D.  
Cloning Distribution: NCI CGAP clone distribution information can be  
found through the I.M.A.G.E. Consortium/ILM at:  
[www-bio.lnlnl.gov/bdnp/image/image.html](http://www-bio.lnlnl.gov/bdnp/image/image.html)  
Insert Length: 770 Std Error: 0.00  
Seq primer: 400p from Gldco.

FEATURES  
SOURCE  
Location/Qualifiers  
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/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="IMAGE:2473948"  
/clone\_lid="NCI CGAP GC6"

BASE COUNT 120 a 95 c 104 t  
ORIGIN  
Query Match 90.3%; Score 336; DB 9; Length 410;  
Best Local Similarity 100.0%; Pred. No. 3.2e-90;  
Matches 336; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Oy 37 GACCTCTTCCGAGAGATTTTATTTATTTATTTATTTATTTATTTATTTATTTAT 96  
Db 18 GACCTCTTCCGAGAGATTTTATTTATTTATTTATTTATTTATTTATTTATTTAT 77  
Oy 97 GAGCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 156  
Db 78 GAGCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 137  
Oy 157 CTGATATCTAGAGAAATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 216  
Db 138 CTGATATCTAGAGAAATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 197  
Oy 217 AATTCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 276  
Db 198 AATTCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 257  
Oy 277 CAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 336  
Db 258 CAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 317  
Oy 337 AACATTTGTCGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 372  
Db 318 AACATTTGTCGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 353

RESULT 3  
A1654552 429 bp mRNA linear EST 17-DEC-1999  
w416b12.x1 NCI CGAP GC6 Homo sapiens CDNA clone IMAGE:230895.3  
DEFINITION  
SIMILAR TO SW:MIK HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM  
CHANNEL PROTEIN ; mRNA sequence.  
ACCESSION  
A1654552.1 GI:4738531  
VERSION  
A1654552  
KEYWORDS  
EST  
ORGANISM  
Homo sapiens  
Human  
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.  
REFERENCE  
NCI CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.  
1 (bases 1 to 429)  
AUTHORS  
National Cancer Institute, Cancer Genome Anatomy Project (CGAP).  
TITLE  
Tumor Gene Index  
JOURNAL  
Contact: Robert Strausberg, Ph.D.  
COMMENT  
Email: [scap@f-mail.nih.gov](mailto:scap@f-mail.nih.gov)  
R. Emmert-Buck, M.D., Ph.D.  
CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima  
Bonaldo, Ph.D.  
DNA Sequencing by: Greg Lennon, Ph.D.  
Cloning Distribution: NCI CGAP clone distribution information can be  
found through the I.M.A.G.E. Consortium/ILM at:  
[www-bio.lnlnl.gov/bdnp/image/image.html](http://www-bio.lnlnl.gov/bdnp/image/image.html)  
Insert Length: 771 Std Error: 0.00

Seq primer: -40BP from Glibco  
High quality sequence stop: 411.  
Location/Qualifiers  
Source

1. 391  
/db.xref="taxon:9606"  
/clone="IMAGE:230895"  
/tissue="NCI-CGAP.GC6"  
/tissue\_type="Pooled germ cell tumors"  
/lab\_host="DH10b"

/note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker. Site 1: Not I; Site 2: Eco RI; plasmid DNA and PCR products were digested with Not I and Eco RI, ligated, and transformed into DH10b cells. The resulting plasmids were isolated and sequenced. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (cloneids 1257096-1258631, 1469064-1470983, and 1475592-1476743). Subtraction by penta Soares and M. Orlina Bonaldio."

BASE COUNT 127 a 100 c 97 g 104 t

Query Match 90.3%; Score 336; DB 9; Length 429;

Best Local Similarity 100.0%; Pred. No. 3.2e-90;

Matches 336; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

37 GAGCTCTCCGAGAGATTATTTACTTATATGAGCAATGGCCCAACACACAGCT 96

12 GAGCTCTCCGAGAGATTATTTACTTATATGAGCAATGGCCCAACACACAGCT 71

97 GAGGAGAGAGCCCTCCACACCAAGATGATGATGAGAGATCTCATATGATCATCTGAC 156

72 GAGGAGAGAGCCCTCCACACCAAGATGATGATGAGAGATCTCATATGATCATCTGAC 131

157 CTCATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 216

132 CTCATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 191

217 AATATCCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 276

192 AATATCCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 251

277 CAGGAAAGATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 336

252 CAGGAAAGATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 311

337 AACATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 372

312 AACATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 347

RESULT 4  
LOCUS A1339609 391 bp mRNA linear EST 29-DEC-1998

DEFINITION g942a07.x1 Soares.NHMPU.SI Homo sapiens cDNA clone IMAGE:1935156

3, similar to SM:MINK.HUMAN P13582 ISK SLOW VOLTAGE-GATED POTASSIUM

CHANNEL PROTEIN ;, mRNA sequence.

ACCESSION A1339609

VERSION A1339609.1

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

1 (bases 1 to 391)

1 (bases 1 to 391)

1 (bases 1 to 391)

1 (bases 1 to 391)

1 (bases 1 to 391)

1 (bases 1 to 391)

1 (bases 1 to 391)

1 (bases 1 to 391)

1 (bases 1 to 391)

Seq primer: -40BP from Glibco  
High quality sequence stop: 380.  
Location/Qualifiers  
Source

1. 391  
/db.xref="taxon:9606"  
/clone="IMAGE:1935156"  
/tissue="NCI-CGAP.NHMPU.SI"  
/tissue\_type="Pooled human melanocyte, fetal heart, and pregnant uterus"

/note="Organ: mixed (see below); Vector: pT73D-Pac (Pharmacia) with a modified polylinker. Site 1: Not I; Site 2: Eco RI; plasmid DNA and PCR products were digested with Not I and Eco RI, ligated, and transformed into DH10b cells. The resulting plasmids were isolated and sequenced. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (cloneids 1257096-1258631, 1469064-1470983, and 1475592-1476743). Subtraction by penta Soares and M. Orlina Bonaldio."

BASE COUNT 119 a 93 c 93 g 86 t

Query Match 89.2%; Score 331.8; DB 9; Length 391;

Best Local Similarity 93.4%; Pred. No. 5.7e-89;

Matches 333; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

38 ACCTCTCCGAGAGATTATTTACTTATATGAGCAATGGCCCAACACACAGCTG 97

3 ACCTCTCCGAGAGATTATTTACTTATATGAGCAATGGCCCAACACACAGCTG 62

98 AGCAGAGAGCCCTCCACACCAAGATGATGATGAGAGATCTCATATGATCATCTGAC 157

63 AGCAGAGAGCCCTCCACACCAAGATGATGATGAGAGATCTCATATGATCATCTGAC 122

158 TCAATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 217

123 TCAATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 182

218 AATCCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 277

183 AATCCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 242

278 AGGAAAGATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 337

243 AGGAAAGATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 302

338 ACATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 372

303 ACATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 337

RESULT 5  
LOCUS A1246239 372 bp mRNA linear EST 28-JAN-1999

DEFINITION g123904.x1 Soares.NHMPU.SI Homo sapiens cDNA clone IMAGE:1857942

3, similar to SM:MINK.HUMAN P13582 ISK SLOW VOLTAGE-GATED POTASSIUM

CHANNEL PROTEIN ;, mRNA sequence.

ACCESSION A1246239

VERSION A1246239.1

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

1 (bases 1 to 372)

1 (bases 1 to 372)

1 (bases 1 to 372)

1 (bases 1 to 372)

1 (bases 1 to 372)

Email: c9apb@fema11.nih.gov  
 This clone is available royalty-free through DLM: contact the  
 National Institutes of Health (NIH) for further information.  
 Insert length: 921 bp  
 Seq primer: 400p from Gdbco  
 High quality sequence stop: 365.

## FEATURES

## SOURCE

Location/Qualifiers

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 /organism="Homo sapiens"  
 /clone="IMAGE:167942"  
 /clone.lib="Searles NIH/PSI"  
 /tissue.type="Pooled human melanocyte, fetal heart, and  
 pregnant uterus"

/lab.host="DH10B"  
 /note="Organ: mixed (see below); Vector: pRTD-Pac  
 Site:1: End:1: Bacterial poly linker; Site:1: Not 1:  
 Site:2: End:1: Bacterial poly linker; Site:2: Not 1:  
 normalized libraries (melanocyte 2NDH, pregnant uterus  
 NBHPV, and fetal heart NBH19V) were mixed, and as circles  
 were made in vitro. Following HAP purification, this DNA  
 was used as tracer in a subtractive hybridization  
 reaction. The driver was PCR-amplified cDNAs from pools of  
 5,000 clones made from the same 3 libraries. The pools  
 340488-345479 and 484488-489479, 9223-26323.

## BASE COUNT

115 a 89 c 86 g 82 t

## ORIGIN

Query Match 88.2%; Score 328; DB 9; Length 372;

## Best Local

Similarity 100.0%; Pred. No. 7/7e-88;

## Matches

328; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

## OR

45 CCGAAGGATTTTATCTATATGACAAATGGCCGACACACAGTGACCAAG 104

## DB

1 CCGAAGGATTTTATCTATATGACAAATGGCCGACACACAGTGACCAAG 60

## OR

105 GGGCCCTCACACCAAGTATGATCTGACAACTCTATATGATCATCTCTACATGAT 164

## DB

61 GGGCCCTCACACCAAGTATGATCTGACAACTCTATATGATCATCTCTACATGAT 120

## OR

165 GATGATGATGATCTCTTTCATCATCGGGCCATCGGTGACACATGTGAATCCAA 224

## DB

121 GATGATGATGATCTCTTTCATCATCGGGCCATCGGTGACACATGTGAATCCAA 180

## OR

225 GAGGCGGCAACATCGAATGACCCCTACACCATCTATGATGAGCATCTGGCAAAA 284

## DB

181 GAGGCGGCAACATCGAATGACCCCTACACCATCTATGATGAGCATCTGGCAAAA 240

## OR

285 GTTACAGAGCCCAATCTGATCTGAGAAAGTCAACAGCCACCATCATGAGAACTGG 344

## DB

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## OR

345 TGGCGGCTGGGTCTAAATGTCTCCCTCA 372

## DB

301 TGGCGGCTGGGTCTAAATGTCTCCCTCA 328

## REPEAT 6

## AK008619

## DEFINITION

## CHANNEL-RELATED PEPTIDE 1 (MIRP) (MIRK-RELATED PEPTIDE 1), full

## ACCESSION

## VERSION

## SOURCE

## ORGANISM

## FEATURES

## source

## location/Qualifiers

## organism="Mus musculus"

## clone="J10R16"

## clone.lib="full-length enriched mouse cDNA library"

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## clone.lib="full-length enriched mouse cDNA library"

## REFERENCE

## AUTHORS

## JOURNAL

## MEDLINE

## PUBMED

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Mammalia: Eutheria, Rodentia, Sciurognathi: Muridae: Murinae: Mus.

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www.bio.11n1.gov/bdip/image/image.html  
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	/clone.type="pooled germ cell tumors"
	/lab.host="DH108"
	/note="Vector: pUT73-Pac (Pharmacia) with a modified
	polylinker: 1st strand cDNA was prepared from 3 pooled
	germ cell tumors, and was then primed with a Not I to Eco
	RI adaptor's (Pharmacia), digested with Not I and cloned
	into the Not I and Eco RI sites of the modified pUT73
	vector. Library is normalized. Library was constructed by
	Bento Soares and M. Fátima Bonaldo."
BASE COUNT	33 a 31 c 28 g 29 t
ORIGIN	
Query Match	33 18; Score 119.4; DB: g; Length 121;
Best Local Similarity 99.28; Percent ID: 25.55;	
Matches 120; Conservative 0; Mismatches 1; Indels 0; Gaps 0;	
Qy 153	GTACCTCATGTGATCATATGAAAGTCTTCATATCATCGGCACTCGTGAGAC 212
Db 1	GTACCTCATGTGATCATATGAAAGTCTTCATATCATCGGCACTCGTGAGAC 60
Qy 213	GTGCAATTCGACAGACCGGACACATCCACATCCACATCCATCATTTATAGCA 272
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Qy 273	C 273
Db 121	C 121
RESULT 13	
LOCUS	AM869303 311 bp mRNA linear EST 22-MAY-2000
DEFINITION	MR3-SN0067-240400-006-f11 SN0067 Homo sapiens cDNA, mRNA sequence.
ACCESSION	AM869303
VERSION	AM869303.1 GI:8003356
KEYWORDS	human.
SOURCE	Human.
ORGANISM	Homo sapiens
REFERENCE	Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
	1 (bases 1 to 311)
	Das Netao, E., Garcia Correa, R., Verjovski-Almeida, S., Briones, M. R.,
	Angel, M. A., de Silva, W. Jr., Zago, R. A., Bordin, S., Costa, F. F.,
	Brundel, L. A., de Oliveira, A. F., Beckner, P., Baldo, S. C., Simpson, D. H.,
	M. J. Soares, F., Brentani, R. R., de Souza, S. J. and
	Simpson, A. J.
TITLE	Shogun sequencing of the human transcriptome with ORF expressed
	sequence tags
JOURNAL	Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
ABSTRACT	20202863 Simpson A. J. G.
COMMENT	Laboratory of Cancer Genetics
	Ludwig Institute for Cancer Research
	Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
	Brazil
	Tel.: +55-11-27049322
	Fax: +55-11-2707001
	Email: asimpson@ludwig.org.br
	This sequence was derived from the RAPD/LIGCH Human Cancer Genome
	Project (http://www.ludwig.org.br/actifs/genhum2.pl?cl=MR3-SN0067-2404
	400-006-ct1c1c-2000-04-24c4a-1)
	Seq primer: puc 18 forward
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source	1. 311 /organism="Homo sapiens" /db="Genbank" /accession="U00007" /def="Homo sapiens" /note="Organ: stomach, normal; Vector: pUC18; Site.1: SmaI; Site.2: SmaI; A mini-library was made by cloning products derived from RESTEST PCR (U.S. Letters patent applications No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC18 vector. Reverse transcription of total RNA was performed using the following conditions: low stringency conditions."
BASE COUNT	60 a 80 c 76 g 95 t
ORIGIN	
Query Match	96.6% Score 99; DB 9; Length 311;
Best Local Similarity	20.6% P: Pred. No. 5.1e-19;
Matches	1; Conservative 10; Mismatches 10; Indels 2; Gaps 1;
Oy	1 ATGCTCACTTTCGCAATTTTCACACAGAGCTGAGACAGCTCTTCGCAACGATTTTAT 60
Db	182 ATGCTCACTTTCGCAATTTTCACACAGAGCTGAGACAGCTCTTCGCAACGATTTTATC 123
Oy	61 ACTTATATGCAATTTGG--CGCCAAACACACAGCTGACGAGAGCCCTCCAGGCCA 118
Db	122 ACTTATATGCAATTTGGCGCGCCGACAGACACAGCTGACGAGAGCCCTCCAGGCCCT 63
Oy	119 AAGTCGATG 127
Db	62 AAGTCGATG 54
RESULT 14	
BB564873	314 bp mRNA linear EST 29-NOV-2000
DEFINITION	BB564873 RIKEN full-length enriched, adult male stomach Mus musculus cDNA clone 220002116 5', mRNA sequence.
VERSION	BB564872
KEYWORDS	BB564873 1 GI:11455765 EST.
SOURCE	house mouse.
ORGANISM	Mus musculus. Strain: B6; Segregation: Chromata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE	1. (bases 1 to 314)
AUTHORS	Alizawa, K., Akahira, S., Altamura, T., Arai, A., Arahata, T., Carninci, P., Hanagaki, K., Hayashi, N., Hirozaki, T., Hirozane, T., Hodojima, Y., Imoh, K., Ishii, Y., Itoh, M., Izawa, K., Kawaji, Y., Kojima, Y., Komine, H., Kusabata, M., Matsuyama, Y., Miyazaki, A., Nakamura, K., Nishi, K., Ohtsuka, K., Shimada, K., Saito, K., Shibata, Y., Shimizu, O., Shibata, Y., Shiraki, T., Sogabe, Y., Suzuki, H., Tagawa, A., Takahashi, F., Tanaka, T., Toyota, T., Methylki, A., Yamanura, T., Yasunishi, A., Yoshida, K., Yoshiki, A., Muramatsu, M. and Hayashizaki, Y.
JOURNAL COMMENT	RIKEN Mouse ESTs (Alizawa, K. et al. 2000) Unpublished (2000) Contract: Yoshida Hayashizaki Laboratory, Genome Research Group, RIKEN Genomic Science Center, 3-1-1 Hirosawa, Yokohama Institute The Institute of Physiological and Chemical Research (RIKEN) 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Fax: 81-45-503-9222 Tel: 81-45-503-9216 Email: genome-res@res.riken.go.jp/ URL: http://genome.res.riken.go.jp/ Carninci, K., Nishiyama, Y., Watanabe, A., Itoh, M., Nagaoaka, S., Sasaki, K., Shimada, K., Hayashizaki, Y., Taniguchi, M., Hayashizaki, Y. Thermoregulation and thermoregulation of thermolabile enzymes by trehalase and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998) Itoh, M., Kitajima, T., Miyama, Y., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki, Y.





Search completed: October 24, 2002, 14:17:22  
Job time : 1610 secs

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GenCore version 5.1.4.P5.4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 20:14:00 ; Search time 21.4496 Seconds

(Without alignments)  
5275.799 Million cell updates/sec

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Scoring table: IDENTITY\_MNC

Gapop 10.0, Gapext 1.0

Searched: 441362 seqs, 153338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database:

Issued Patents NA.\*

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5: /cgn2.6/product1/lna/PCFUS.COMB.seq.\*  
6: /cgn2.6/product1/lna/Dockfilist.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No	Query Score	Match Length	DB ID	Description
1	53.2	14.4	398 1	US-08-118-101A-5
2	53.2	14.4	436 6	US-09-679-185-1
3	53.2	14.4	1703 3	US-09-135-021-77
4	53.2	14.4	1703 4	US-09-135-020-3
5	53.2	14.4	1703 4	US-09-135-010A-3
6	53.2	14.4	1703 4	US-09-597-735-3
7	53.2	14.4	1703 4	US-09-444-295-3
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11	36.6	9.9	2652 1	US-08-110-286A-1
12	36.6	9.9	1880 1	US-08-462-746-1
13	36.6	9.9	1880 1	US-08-462-746-1
14	36.6	9.9	1880 1	US-08-462-746-1
15	33.6	9.1	645 3	US-09-069-896-2
16	33.6	9.1	645 3	US-09-471-466-2
17	33.6	8.9	606 4	US-09-328-111-133
18	33.6	8.9	2912 4	US-09-307-143-3
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#### ALIGNMENTS

RESULT 1  
US-08-118-101A-5  
; Sequence 5, Application US/08118101A  
; Patent No. 5620892  
; GENERAL INFORMATION:  
; INVENTOR: KUTZ, Stephen E.  
; APPLICANT: Bristol-Myers Squibb, LLC  
; TITLE OF INVENTION: A STRAIN OF SACHAROWYCES CEREVISIAE  
; NUMBER OF SEQUENCES: 16  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Burton Rodney  
; CITY: Princeton  
; STATE: New Jersey  
; ZIP: 08543-4000  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; CONVERTER: IBM PC compatible  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/118,101A  
; FILING DATE:  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: CAUL, Timothy J.  
; REFERENCE/DOCKET NUMBER: DC27  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (609) 252-5901  
; INFORMATION FOR SEQ ID NO: 5:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 358 bases  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: 1..398  
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Best Local Similarity 62.6%; Pred. No. 1.7e+08;  
Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;



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FILE REFERENCE: 2323-131
CURRENT APPLICATION NUMBER: US/09/444,871
EARLIER FILING DATE: 1999-11-22
EARLIER APPLICATION NUMBER: 09/135,020
EARLIER FILING DATE: 1998-08-29
EARLIER APPLICATION NUMBER: 08/921,066
EARLIER FILING DATE: 1997-08-29
EARLIER APPLICATION NUMBER: 08/739,383
EARLIER FILING DATE: 1996-10-29
EARLIER APPLICATION NUMBER: 06/019,014
EARLIER FILING DATE: 1995-12-01
EARLIER APPLICATION NUMBER: 60/094,477
NUMBER OF SEQ ID NOS: 114
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 3
LENGTH: 1703
TYPE: DNA
ORGANISM: Homo sapiens
FEATURES:
LOCATION: CDS
LOCATION: (135)..(579)
US-09-444-871-3

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Db      324 CCTGACGCTCTCATGCGTAGTGAGATTCTTGCGCTTCTCACACCCTTGCCATGCGTGG
Qy      210 CACTGTGATTCACAAAGACGGGAACCTCCATACCCCTTACACAGTAATTC--T 266
Db      364 CTGACGTCGCTCCAAAGATCGAAGACGACGACCATTCACAGCTACATCATGATGC 443
Qy      267 AGTACGTCGCGGAGAAGATCGAACGCGCAATC 300
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RESULT 7
Sequence 3, Application US/09597735
US-09-597-735-3
SEQUENCE: 6401130
GENERAL INFORMATION:
APPLICANT: Keating, Mark T.
APPLICANT: Sanguinetti, Michael C.
APPLICANT: Curran, Mark E.
APPLICANT: Landes, Gregory M.
APPLICANT: Comors, Timothy D.
APPLICANT: Burn, Timothy C.
APPENDIX: SEQUENCES
TITLE OF INVENTION: KLOTZ - A LONG QT SYNDROME GENE
FILE REFERENCE: 2323-133
CURRENT APPLICATION NUMBER: US/09/597,735
CURRENT FILING DATE: 2000-06-19
EARLIER APPLICATION NUMBER: 09/135,010
EARLIER FILING DATE: 1998-08-17
EARLIER APPLICATION NUMBER: 60/094,477
EARLIER FILING DATE: 1998-08-29
EARLIER APPLICATION NUMBER: 08/921,066
EARLIER FILING DATE: 1997-08-29
EARLIER APPLICATION NUMBER: 08/739,383
EARLIER FILING DATE: 1996-10-29
EARLIER APPLICATION NUMBER: 60/019,014
EARLIER FILING DATE: 1995-12-22
NUMBER OF SEQ ID NOS: 116
SOFTWARE: PatentIn ver. 2.0
SEQ ID NO 3
LENGTH: 1703
TYPE: DNA
ORGANISM: Homo sapiens

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LOCATION: 82, 1329  
 OTHER INFORMATION: /product= "HUMAN PITUITARY"  
 OTHER INFORMATION: CRF-RECEPTOR\*  
 OTHER INFORMATION: /note= "this sequence is encoded by clone  
 OTHER INFORMATION: "CRF-R1".\*  
 US-08-110-286A-1

Query Match 9.8%; Score 36; DB 1; Length 180;  
 Best Local Similarity 52.7%; Pred. No. 0.025; Indels 0; Gaps 0;

Matches 78; Conservative 0; Mismatches 70; Indels 0; Gaps 0;

OY 115 GCCAAGTGTACTGAGAACTTCTACATGCTGCTGATCGTATGATGGA 174  
 DB 862 GGCAGAAAGCCCTGGGGTGTACACGACATCACTACAGGCCCATGATCTGGTCTG 921  
 OY 115 ATGTCCTTCATCATCGTGGCCATCGGTGAGCACTGTAAATTCAGAGCGGAA 234  
 DB 932 CTGATCATTTTCATCTCTTTCATCATGCTGCGCATCCCTATGACCAAGCTCGGGCA 981  
 OY 235 CACTGCATATGACCCCTACACCACTGACA 262  
 DB 982 TCCACACGCTGTACACATTCAGTACA 1009

RESULT 13  
 US-08-482-746-1

/sequence 1, Application US/08482746B  
 / Patent No. 6399315

/ GENERAL INFORMATION:  
 / APPLICANT: Pettin, Marilyn H.  
 / APPLICANT: Chen, Rongping  
 / APPLICANT: Lewis, Kathy A.  
 / APPLICANT: Vale Jr., Wylie W.  
 / APPLICANT: Donaldson, Cynthia J.  
 / APPLICANT: Sarmiento, Paul  
 / TITLE OF INVENTION: Cloning and Recombinant Production of  
 / TITLE OF INVENTION: CRF Receptor(s)  
 / FILE REFERENCE: P41-90002

/ CURRENT FILING DATE: 1995-06-07  
 / EARLIER FILING DATE: 1995-01-17  
 / EARLIER APPLICATION NUMBER: US 08/353,537  
 / EARLIER FILING DATE: 1994-05-12  
 / EARLIER APPLICATION NUMBER: US 08/353,537  
 / EARLIER FILING DATE: 1994-05-25  
 / EARLIER APPLICATION NUMBER: US 08/110,286  
 / EARLIER FILING DATE: 1993-08-23  
 / EARLIER APPLICATION NUMBER: US 08/079,320  
 / NUMBER OF SEQ NOS: 13  
 / SOFTWARE: FASTSEQ for Windows Version 3.0

/ SEQ ID NO 1  
 / LENGTH: 1495  
 / TYPE: DNA  
 / ORGANISM: Homo sapiens  
 / FEATURE:  
 / NAME/REV: CDS  
 / LOCATION: (82)...(1326)  
 / OTHER INFORMATION: /product= "Human pituitary CRF-receptor"  
 / OTHER INFORMATION: /note= "this sequence is encoded by clone  
 / OTHER INFORMATION: "CRF-R1".\*  
 US-08-482-746-1

Query Match 9.8%; Score 36; DB 4; Length 1495;  
 Best Local Similarity 52.7%; Pred. No. 0.026;

Matches 78; Conservative 0; Mismatches 70; Indels 0; Gaps 0;

OY 115 GCCAAGTGTACTGAGAACTTCTACATGCTGCTGATCGTATGATGGA 174  
 DB 862 GGCAGAAAGCCCTGGGGTGTACACGACATCACTACAGGCCCATGATCTGGTCTG 921  
 OY 175 ATGTCCTTCATCATCGTGGCCATCGGTGAGCACTGTAAATTCAGAGCGGAA 234

DB 922 CTGATCATTTTCATCTCTTTCATCATGCTGCGCATCCCTATGACCAAGCTCGGGCA 981  
 OY 235 CACTGCATATGACCCCTACACCACTGACA 262  
 DB 982 TCCACACGCTGTACACATTCAGTACA 1009

RESULT 14  
 US-08-482-746-14

/sequence 14, Application US/08482746B  
 / Patent No. 6399315

/ GENERAL INFORMATION:  
 / APPLICANT: Pettin, Marilyn H.  
 / APPLICANT: Chen, Rongping  
 / APPLICANT: Lewis, Kathy A.  
 / APPLICANT: Vale Jr., Wylie W.  
 / APPLICANT: Donaldson, Cynthia J.  
 / APPLICANT: Sarmiento, Paul  
 / TITLE OF INVENTION: Cloning and Recombinant Production of  
 / TITLE OF INVENTION: CRF Receptor(s)  
 / FILE REFERENCE: P41-90002

/ CURRENT FILING DATE: 1995-06-07  
 / EARLIER FILING DATE: 1995-01-17  
 / EARLIER APPLICATION NUMBER: US 08/353,537  
 / EARLIER FILING DATE: 1994-05-12  
 / EARLIER APPLICATION NUMBER: US 08/353,537  
 / EARLIER FILING DATE: 1994-05-25  
 / EARLIER APPLICATION NUMBER: US 08/110,286  
 / EARLIER FILING DATE: 1993-08-23  
 / EARLIER APPLICATION NUMBER: US 08/079,320  
 / NUMBER OF SEQ ID NOS: 13  
 / SOFTWARE: FASTSEQ for Windows Version 3.0

/ SEQ ID NO 14  
 / LENGTH: 1582  
 / TYPE: DNA  
 / ORGANISM: Homo sapiens  
 / FEATURE:  
 / NAME/REV: CDS  
 / LOCATION: (82)...(1413)  
 / OTHER INFORMATION: CRF-R splice-variant Insert fragment inserted  
 / OTHER INFORMATION: between nucleotides 516-517 of SEQ ID NO:1.  
 / OTHER INFORMATION: /note= "this sequence is contained in clone  
 / OTHER INFORMATION: "CRF-R2".\*  
 US-08-482-746-14

Query Match 9.8%; Score 36; DB 4; Length 1582;  
 Best Local Similarity 52.7%; Pred. No. 0.027;

Matches 78; Conservative 0; Mismatches 70; Indels 0; Gaps 0;

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 OY 175 ATGTCCTTCATCATCGTGGCCATCGGTGAGCACTGTAAATTCAGAGCGGAA 234  
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 OY 235 CACTGCATATGACCCCTACACCACTGACA 262  
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RESULT 15  
 US-09-069-896-2

/sequence 2, Application US/0906986  
 / Patent No. 6071720

/ GENERAL INFORMATION:  
 / APPLICANT: Pettin, Marilyn H.  
 / APPLICANT: Patterson, Chandra



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? APPLICANT: Corley, Neil C.
? TITLE OF INVENTION: DELTAED RECTIFIER POTASSIUM
? TITLE OF INVENTION: CHANNEL HOMOLOG
? NUMBER OF SEQUENCES: 4
? CORRESPONDENCE ADDRESS:
? ADDRESS: Indyte Pharmaceuticals, Inc.
? STREET: 3174 Rafter Drive
? CITY: Palo Alto
? STATE: CA
? COUNTRY: USA
? ZIP: 94304
? COMPUTER READABLE FORM:
? MODIFIED: 05/09/96
? MODIFIED BY: JRM
? OPERATING SYSTEM: DOS
? SOFTWARE: FASTSEQ for Windows Version 2.0
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: US/09/069,896
? FILING DATE:
? CLASSIFICATION:
? PRIOR APPLICATION DATA:
? APPLICATION NUMBER:
? ATTORNEY/AGENT INFORMATION:
? NAME: Ceitone, Michael C
? REGISTRATION NUMBER: 39,132
? TELECOMMUNICATION INFORMATION:
? TELEPHONE: 650 853-0555
? TELEFAX: 650 845-4166
? TELEX:
? INFORMATION FOR SEQ ID NO: 2:
? SEQUENCE CHARACTERISTICS:
? LENGTH: 645 base pairs
? TYPE: nucleic acid
? STRANDEDNESS: single
? ORIENTATION: forward
? IMMEDIATE SOURCE:
? LIBRARY: BRSTNOT03
? CLONE: 637471
? US-09-069-896-2

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Query Match          5 18; Score 33.6; DB 3; Length 645;
Similarity          53 96; Percent 0.1;
Matches          69; Conservative 0; Mismatches 59; Indels 0; Gaps 0;

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Db 320 GGAAGCCGC 327

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